

# WEST Search History

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DATE: Thursday, September 09, 2004

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<input type="checkbox"/>	L19	L15 AND AScr	27
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<input type="checkbox"/>	L12	L11 AND PrP	40
<input type="checkbox"/>	L11	L10 AND prion	273
<input type="checkbox"/>	L10	514/2,4,12.CCLS.	12236
<input type="checkbox"/>	L9	L8 AND PrPSc	3
<input type="checkbox"/>	L8	L7 AND PrP	20
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**Search Results - Record(s) 1 through 27 of 27 returned.** 1. Document ID: US 20040147531 A1**Using default format because multiple data bases are involved.**

L19: Entry 1 of 27

File: PGPB

Jul 29, 2004

PGPUB-DOCUMENT-NUMBER: 20040147531

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040147531 A1

TITLE: Amidine derivatives for treating amyloidosis

PUBLICATION-DATE: July 29, 2004

**INVENTOR-INFORMATION:**

NAME	CITY	STATE	COUNTRY	RULE-47
Chalifour, Robert J.	Ille Bizard		CA	
Kong, Xianqi	Pierrefonds		CA	
Wu, Xinfu	Dollard-des-Ormeaux		CA	
Lu, Wenshuo	LaSalle		CA	

US-CL-CURRENT: [514/256](#); [514/397](#), [514/636](#)

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KMC](#) | [Drawn Desc](#)

 2. Document ID: US 20040138178 A1

L19: Entry 2 of 27

File: PGPB

Jul 15, 2004

PGPUB-DOCUMENT-NUMBER: 20040138178

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040138178 A1

TITLE: Phosphono-carboxylate compounds for treating amyloidosis

PUBLICATION-DATE: July 15, 2004

**INVENTOR-INFORMATION:**

NAME	CITY	STATE	COUNTRY	RULE-47
Szarek, Walter A.	Kingston		CA	
Kong, Xianqi	Pierrefonds		CA	
Thatcher, Gregory R.J.	Kingston		CA	
Gorine, Boris	Edmonton		CA	

US-CL-CURRENT: [514/79](#); [514/114](#), [514/141](#)**ABSTRACT:**

h e b b g e e e f e b ef b e

Therapeutic compounds and methods for modulating amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is modulated by the administration to a subject of an effective amount of a therapeutic compound comprising a phosphonate group and a carboxylate group, a congener thereof, or a pharmaceutically acceptable salt or ester thereof. In preferred embodiments, an interaction between an amyloidogenic protein and a basement membrane constituent is modulated.

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KMC](#) | [Draw. Desc](#)

3. Document ID: US 20040048279 A1

L19: Entry 3 of 27

File: PGPB

Mar 11, 2004

PGPUB-DOCUMENT-NUMBER: 20040048279  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20040048279 A1

TITLE: Method for detecting methylation states for a toxicological diagnostic

PUBLICATION-DATE: March 11, 2004

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Olek, Alexander	Berlin		DE	
Piepenbrock, Christian	Berlin		DE	
Berlin, Kurt	Stahnsdorf		DE	

US-CL-CURRENT: 435/6

ABSTRACT:

The present invention concerns a method for toxicological diagnosis. A DNA sample is taken from an organism or a cell culture, which has previously been subjected to a specific substance that is to be investigated for its toxicological effect. The DNA contained in this sample is chemically pretreated and the base sequence of a part of the modified DNA is determined. A methylation state characteristic for the sample or a characteristic methylation pattern is concluded from this. The effect of a substance on the organism or the cell culture is concluded by comparison with data of the methylation states of other samples and/or compared with other substances from a toxicological point of view.

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KMC](#) | [Draw. Desc](#)

4. Document ID: US 20040006092 A1

L19: Entry 4 of 27

File: PGPB

Jan 8, 2004

PGPUB-DOCUMENT-NUMBER: 20040006092  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20040006092 A1

TITLE: Amidine derivatives for treating amyloidosis

h e b b g e e e f e b ef b e

PUBLICATION-DATE: January 8, 2004

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Chalifour, Robert J.	Ile Bizard		CA	
Kong, Xianqi	Dollard-des-Ormeaux		CA	
Wu, Xinfu	Dollard-des-Ormeaux		CA	
Lu, Wenshuo	Montreal		CA	

US-CL-CURRENT: 514/256; 514/397, 514/632

## ABSTRACT:

The present invention relates to the use of amidine compounds in the treatment of amyloid-related diseases. In particular, the invention relates to a method of treating or preventing an amyloid-related disease in a subject comprising administering to the subject a therapeutic amount of an amidine compound. Among the compounds for use according to the invention are those according to the following Formula, such that, when administered, amyloid fibril formation, neurodegeneration, or cellular toxicity is reduced or inhibited: 1

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KMC](#) | [Drawn Desc](#)

5. Document ID: US 20030236392 A1

L19: Entry 5 of 27

File: PGPB

Dec 25, 2003

PGPUB-DOCUMENT-NUMBER: 20030236392

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030236392 A1

TITLE: Novel full length cDNA

PUBLICATION-DATE: December 25, 2003

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Isogai, Takao	Ibaraki		JP	
Sugiyama, Tomoyasu	Tokyo		JP	
Otsuki, Tetsuji	Chiba		JP	
Wakamatsu, Ai	Chiba		JP	
Sato, Hiroyuki	Osaka		JP	
Ishii, Shizuko	Chiba		JP	
Yamamoto, Jun-ichi	Chiba		JP	
Isono, Yuuko	Chiba		JP	
Hio, Yuri	Chiba		JP	
Otsuka, Kaoru	Saitama		JP	
Nagai, Keiichi	Tokyo		JP	
Irie, Ryotaro	Chiba		JP	
Tamechika, Ichiro	Osaka		JP	
Seki, Naohiko	Chiba		JP	
Yoshikawa, Tsutomu	Chiba		JP	
Otsuka, Motoyuki	Tokyo		JP	

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Nagahari, Kenji	Tokyo	JP
Masuho, Yasuhiko	Tokyo	JP

US-CL-CURRENT: 536/23.1; 435/183, 435/325, 435/6, 435/69.1, 530/350, 702/19

## ABSTRACT:

Novel full-length cDNAs are provided.

1970 cDNA derived from human have been isolated. The full-length nucleotide sequences of the cDNA and amino acid sequences encoded by the nucleotide sequences have been determined. Because the cDNA of the present invention are full-length and contain the translation start site, they provide information useful for analyzing the functions of the polypeptide.

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KWC](#) | [Drawn Des](#)

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**6. Document ID: US 20030232758 A1**

L19: Entry 6 of 27

File: PGPB

Dec 18, 2003

PGPUB-DOCUMENT-NUMBER: 20030232758

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030232758 A1

TITLE: Immunological methods and compositions for the treatment of Alzheimer's disease

PUBLICATION-DATE: December 18, 2003

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
St. George-Hyslop, Peter H.	Toronto		CA	
McLaurin, JoAnne	Toronto		CA	

US-CL-CURRENT: 514/12; 435/320.1, 435/325, 435/69.1, 530/324, 536/23.1

## ABSTRACT:

The present invention relates to immunogenic compositions and peptides comprising residues 4-10 (FRHDSGY) of the amyloid peptide Abeta<sub>sub.42</sub>. The invention further relates to antibodies that bind to the Abeta<sub>sub.(4-10)</sub> antigenic determinant. The invention provides methods for treating Alzheimer's disease and for reducing the amyloid load in Alzheimers patients. The invention also relates to methods for designing small molecule inhibitors of amyloid deposition.

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KWC](#) | [Drawn Des](#)

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**7. Document ID: US 20030185808 A1**

L19: Entry 7 of 27

File: PGPB

Oct 2, 2003

PGPUB-DOCUMENT-NUMBER: 20030185808

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PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20030185808 A1

TITLE: Prostate cell lines

PUBLICATION-DATE: October 2, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Thraves, Peter	London		GB	
Sutton, Andrew	London		GB	

US-CL-CURRENT: 424/93.21; 424/85.2, 435/366, 514/44

ABSTRACT:

An increasingly aged population and better diagnosis has lead to an apparent increase in the prevalence of prostate cancer in men. There is an acute need to better understand the progression of this disease from its locally confined site of initiation to the end stage widely metastatic disease with attendant morbidity and mortality. It has historically been difficult to raise and maintain immortalised prostate cell lines in culture. We have derived a cell line selected from the group consisting of clones ONYCAP 1 and ONYCAP23. The cell lines are characterised as being prostate epithelial in origin.

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KOMC](#) | [Drawn Desc](#)

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8. Document ID: US 20030108595 A1

L19: Entry 8 of 27

File: PGPB

Jun 12, 2003

PGPUB-DOCUMENT-NUMBER: 20030108595  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20030108595 A1

TITLE: Method for treating amyloidosis

PUBLICATION-DATE: June 12, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Kisilevsky, Robert	Kingston		CA	
Szarek, Walter	Kingston		CA	
Weaver, Donald	Kingston		CA	

US-CL-CURRENT: 424/450; 514/12, 514/23, 514/378, 514/381, 514/460, 514/79

ABSTRACT:

Therapeutic compounds and methods for inhibiting amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is inhibited by the administration to a subject of an effective amount of a therapeutic compound comprising an anionic group and a carrier molecule, or a pharmaceutically acceptable salt thereof, such that an interaction between an amyloidogenic protein and a basement membrane constituent is inhibited. Preferred anionic groups are sulfonates

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and sulfates. Preferred carrier molecules include carbohydrates, polymers, peptides, peptide derivatives, aliphatic groups, alicyclic groups, heterocyclic groups, aromatic groups and combinations thereof.

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KMC](#) | [Draw. Des.](#)

9. Document ID: US 20030027796 A1

L19: Entry 9 of 27

File: PGPB

Feb 6, 2003

PGPUB-DOCUMENT-NUMBER: 20030027796

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030027796 A1

TITLE: Phosphono-carboxylate compounds for treating amyloidosis

PUBLICATION-DATE: February 6, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Szarek, Walter A.	Kingston		CA	
Kong, Xianqi	Dollard-des-Ormeaux		CA	
Thatcher, Gregory R.J.	Kingston		CA	
Gorine, Boris	Edmonton		CA	

US-CL-CURRENT: 514/79; 514/114, 514/141

ABSTRACT:

Therapeutic compounds and methods for modulating amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is modulated by the administration to a subject of an effective amount of a therapeutic compound comprising a phosphonate group and a carboxylate group, a congener thereof, or a pharmaceutically acceptable salt or ester thereof. In preferred embodiments, an interaction between an amyloidogenic protein and a basement membrane constituent is modulated.

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KMC](#) | [Draw. Des.](#)

10. Document ID: US 20020119926 A1

L19: Entry 10 of 27

File: PGPB

Aug 29, 2002

PGPUB-DOCUMENT-NUMBER: 20020119926

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020119926 A1

TITLE: Inhibitors of IAPP fibril formation and uses thereof

PUBLICATION-DATE: August 29, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
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e b

ef b e

Fraser, Paul

Toronto

CA

US-CL-CURRENT: 514/12; 435/184, 514/14, 514/15, 514/16, 514/17

## ABSTRACT:

New antifibrilllogenic agents and compositions containing same, methods of using the antifibrilllogenic agents and compositions for inhibiting amyloid fibril formation, and effective therapeutics for preventing or delaying the progression of, e.g., Alzheimer's disease and diabetes.

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KMC | Drawn Desc

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11. Document ID: US 20020115717 A1

L19: Entry 11 of 27

File: PGPB

Aug 22, 2002

PGPUB-DOCUMENT-NUMBER: 20020115717

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020115717 A1

TITLE: Amyloid targeting imaging agents and uses thereof

PUBLICATION-DATE: August 22, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Gervais, Francine	Ile Bizard		CA	
Kong, Xianqi	Dollard-des-Ormeaux		CA	
Chalifour, Robert	Ile Bizard		CA	
Migneault, David	Laval		CA	

US-CL-CURRENT: 514/553; 424/1.11

## ABSTRACT:

Amyloid-targeting imaging agents such as radiolabeled amyloid targeting molecules and amyloid targeting molecule-chelator conjugates for imaging, e.g., amyloid plaques in vivo, and/or for the treatment of amyloidosis disorders. The invention provides amyloid-targeting imaging agents that are useful for imaging sites of amyloid disease. Imaging agents of the invention are capable of binding specifically to amyloid plaques, as an aid in diagnosis and/or early treatment of amyloidosis disorders.

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KMC | Drawn Desc

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12. Document ID: US 20020094335 A1

L19: Entry 12 of 27

File: PGPB

Jul 18, 2002

PGPUB-DOCUMENT-NUMBER: 20020094335

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020094335 A1

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TITLE: Vaccine for the prevention and treatment of alzheimer's and amyloid related diseases

PUBLICATION-DATE: July 18, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Chalifour, Robert	Ile Bizard		CA	
Hebert, Lise	Brossard		CA	
Kong, Xianqi	Dollard-des-Oremaux		CA	
Gervais, Francine	Ile Bizard		CA	

US-CL-CURRENT: 424/185.1

ABSTRACT:

The present invention relates to a stereochemically based "non-self" antigen vaccine for the prevention and/or treatment of Alzheimer's and other amyloid related diseases. The present invention provides a vaccine for the prevention and treatment of Alzheimer's and other amyloid related diseases, which overcomes the drawbacks associated with using naturally occurring peptides, proteins or immunogens.

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KMC](#) | [Draw Desc](#)

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13. Document ID: US 20020009730 A1

L19: Entry 13 of 27

File: PGPB

Jan 24, 2002

PGPUB-DOCUMENT-NUMBER: 20020009730

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020009730 A1

TITLE: Human stress array

PUBLICATION-DATE: January 24, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Chenchik, Alex	Palo Alto	CA	US	
Lukashev, Matvey E.	Newton	MA	US	

US-CL-CURRENT: 435/6; 536/24.3

ABSTRACT:

Human stress arrays and methods for their use are provided. The subject arrays include a plurality of polynucleotide spots, each of which is made up of a polynucleotide probe composition of unique polynucleotides corresponding to a human stress gene. The subject arrays find use in hybridization assays, particularly in assays for the identification of differential gene expression of human stress genes.

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KMC](#) | [Draw Desc](#)

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## 14. Document ID: US 20010048941 A1

L19: Entry 14 of 27

File: PGPB

Dec 6, 2001

PGPUB-DOCUMENT-NUMBER: 20010048941  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20010048941 A1

TITLE: Method for treating amyloidosis

PUBLICATION-DATE: December 6, 2001

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Kisilevsky, Robert	Kingston		CA	
Szarek, Walter	Kingston		CA	
Weaver, Donald	Kingston		CA	

US-CL-CURRENT: 424/450; 514/2, 514/378, 514/381, 514/460, 514/54

## ABSTRACT:

Therapeutic compounds and methods for inhibiting amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is inhibited by the administration to a subject of an effective amount of a therapeutic compound comprising an anionic group and a carrier molecule, or a pharmaceutically acceptable salt thereof, such that an interaction between an amyloidogenic protein and a basement membrane constituent is inhibited. Preferred anionic groups are sulfonates and sulfates. Preferred carrier molecules include carbohydrates, polymers, peptides, peptide derivatives, aliphatic groups, alicyclic groups, heterocyclic groups, aromatic groups and combinations thereof

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KMC](#) | [Draw. Desc](#)

## 15. Document ID: US 20010027186 A1

L19: Entry 15 of 27

File: PGPB

Oct 4, 2001

PGPUB-DOCUMENT-NUMBER: 20010027186  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20010027186 A1

TITLE: Phosphono-carboxylate compounds for treating amyloidosis

PUBLICATION-DATE: October 4, 2001

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Szarek, Walter A.	Kingston		CA	
Kong, Xianqi	Dollard-des-Ormeaux		CA	
Thatcher, Gregory R.J	Kingston		CA	
Gorine, Boris	Edmonton		CA	

US-CL-CURRENT: 514/79; 514/114, 514/129, 514/142

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## ABSTRACT:

Therapeutic compounds and methods for modulating amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is modulated by the administration to a subject of an effective amount of a therapeutic compound comprising a phosphonate group and a carboxylate group, a congener thereof, or a pharmaceutically acceptable salt or ester thereof. In preferred embodiments, an interaction between an amyloidogenic protein and a basement membrane constituent is modulated.

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KOMC](#) | [Draw. Desc](#)

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□ 16. Document ID: US 6632808 B1

L19: Entry 16 of 27

File: USPT

Oct 14, 2003

US-PAT-NO: 6632808

DOCUMENT-IDENTIFIER: US 6632808 B1

TITLE: Inhibitors of amyloid formation

DATE-ISSUED: October 14, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Caughey; Winslow S.	Hamilton	MT		
Caughey; Byron	Hamilton	MT		

US-CL-CURRENT: 514/185, 514/410, 540/122, 540/145

ABSTRACT:

Methods, compounds and compositions are disclosed for treating amyloidogenic diseases, like Alzheimer's disease and type 2 diabetes, and particularly prion diseases associated with conversion of protease sensitive PrP (PrP-sen) to protease resistant PrP (PrP-res), by administering therapeutically effective amounts of a tetrapyrrole. Particular disclosed tetrapyrroles having this activity include phthalocyanines, deuteroporphyrins, and meso-substituted porphines. Complexes of certain of the pyrroles with metals or metal ions produce compounds that are particularly effective in converting the conversion of PrP-sen to PrP-res. The treatment of the present invention is particularly suited for preventing or inhibiting the progression of prion related diseases, such as transmissible spongiform encephalopathies.

70 Claims, 10 Drawing figures

Exemplary Claim Number: 1

Number of Drawing Sheets: 4

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KOMC](#) | [Draw. Desc](#)

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□ 17. Document ID: US 6562836 B1

L19: Entry 17 of 27

File: USPT

May 13, 2003

US-PAT-NO: 6562836

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DOCUMENT-IDENTIFIER: US 6562836 B1

\*\* See image for Certificate of Correction \*\*

TITLE: Methods and compounds for inhibiting amyloid deposits

DATE-ISSUED: May 13, 2003

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Szarek; Walter A.	Kingston			CA
Weaver; Donald F.	Kingston			CA
Kong; Xianqi	Dollard-des-Ormeaux			CA
Gupta; Ajay	Pointe-Claire			CA
Migneault; David	Laval			CA

US-CL-CURRENT: 514/307; 514/308, 514/311, 514/313, 514/314

## ABSTRACT:

Methods and compositions which are useful in the treatment of amyloidosis. In particular, methods and compositions are provided for inhibiting, preventing and treating amyloid deposition, e.g., in pancreatic islets, wherein the amyloidotic deposits are islet amyloid polypeptide (IAPP)-associated amyloid deposition or deposits. The methods of the invention involve administering to a subject a therapeutic compound which inhibits IAPP-associated amyloid deposits. Accordingly, the compositions and methods of the invention are useful for inhibiting IAPP-associated amyloidosis in disorders in which such amyloid deposition occurs, such as diabetes.

172 Claims, 14 Drawing figures

Exemplary Claim Number: 1

Number of Drawing Sheets: 14

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | | | [Claims](#) | [KIMC](#) | [Draw. Desc.](#)

## 18. Document ID: US 6440952 B2

L19: Entry 18 of 27

File: USPT

Aug 27, 2002

US-PAT-NO: 6440952

DOCUMENT-IDENTIFIER: US 6440952 B2

TITLE: Phosphono-carboxylate compounds for treating amyloidosis

DATE-ISSUED: August 27, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Szarek; Walter A.	Kingston			CA
Kong; Xianqi	Dollard-des-Ormeaux			CA
Thatcher; Gregory R. J.	Kingston			CA
Gorine; Boris	Edmonton			CA

US-CL-CURRENT: 514/120; 558/110, 558/70

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## ABSTRACT:

Therapeutic compounds and methods for modulating amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is modulated by the administration to a subject of an effective amount of a therapeutic compound comprising a phosphonate group and a carboxylate group, a congener thereof, or a pharmaceutically acceptable salt or ester thereof. In preferred embodiments, an interaction between an amyloidogenic protein and a basement membrane constituent is modulated.

20 Claims, 0 Drawing figures

Exemplary Claim Number: 1

Full Title Citation Front Review Classification Date Reference     Claims IOMC Drawn Desc

19. Document ID: US 6355784 B1

L19: Entry 19 of 27

File: USPT

Mar 12, 2002

US-PAT-NO: 6355784

DOCUMENT-IDENTIFIER: US 6355784 B1

**\*\* See image for Certificate of Correction \*\***

TITLE: Methods and compositions for the manufacture of halogenated anthracyclines with increased antitumor activity, other anthracyclines, halogenated sugars, and glycosyl donors

DATE-ISSUED: March 12, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Priebe; Waldemar	Houston	TX	77005	
Krawczyk; Marta	Lexington	KY	40503	
Skibicki; Piotr	Warsaw 04015			PL
Fokt; Izabela	The Woodlands	TX	77380	
Dziewiszek; Krzysztof	The Woodlands	TX	77380	
Grynkiewicz; Grzegorz	05-092 Lomianki			PL
Perez-Soler; Roman	New York	NY	10016	

US-CL-CURRENT: 536/6.4; 536/122, 536/17.2, 536/18.4, 536/18.7, 536/4.1

ABSTRACT:

The present invention discloses new and novel halogenated anthracyclines linked through the saccharide portions. These congeners show high activity in vitro against several tumor cell lines. In doxorubicin (DOX) sensitive cell lines, they are at least as cytotoxic as DOX and in some cases more so. Many of these 4'- and 6'-fluorinated anthracyclines are more effective against multidrug-resistant tumors than was DOX, and/or have greater effectiveness than DOX against DOX sensitive cells. The compounds of this invention also have anti-amyloidogenic effects and the use of these compounds in the treatment of Alzheimer's disease is contemplated.

7 Claims, 19 Drawing figures

Exemplary Claim Number: 1

Number of Drawing Sheets: 15

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Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KOMC	Drawn Des.
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20. Document ID: US 6329356 B1

L19: Entry 20 of 27

File: USPT

Dec 11, 2001

US-PAT-NO: 6329356

DOCUMENT-IDENTIFIER: US 6329356 B1

TITLE: Phosphono-carboxylate compounds for treating amyloidosis

DATE-ISSUED: December 11, 2001

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Szarek; Walter A.	Kingston			CA
Kong; Xianqi	Dollard-des-Ormeaux			CA

US-CL-CURRENT: 514/120

## ABSTRACT:

Therapeutic compounds and methods for modulating amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is modulated by the administration to a subject of an effective amount of a therapeutic compound comprising a phosphonate group and a carboxylate group, a congener thereof, or a pharmaceutically acceptable salt or ester thereof. In preferred embodiments, an interaction between an amyloidogenic protein and a basement membrane constituent is modulated.

31 Claims, 0 Drawing figures

Exemplary Claim Number: 1

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KOMC	Drawn Des.
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21. Document ID: US 5972328 A

L19: Entry 21 of 27

File: USPT

Oct 26, 1999

US-PAT-NO: 5972328

DOCUMENT-IDENTIFIER: US 5972328 A

\*\* See image for Certificate of Correction \*\*

TITLE: Method for treating amyloidosis

DATE-ISSUED: October 26, 1999

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Kisilevsky; Robert	Kingston			CA
Szarek; Walter	Kingston			CA
Weaver; Donald	Kingston			CA

US-CL-CURRENT: 424/78.31; 424/423, 424/427, 424/430, 424/434, 424/436, 424/441,

h e b b g e e e f e b ef b e

424/450, 424/78.35, 514/772.4, 526/286, 526/287

## ABSTRACT:

Therapeutic compounds and methods for inhibiting amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is inhibited by the administration to a subject of an effective amount of a therapeutic compound comprising an anionic group and a carrier molecule, or a pharmaceutically acceptable salt thereof, such that an interaction between an amyloidogenic protein and a basement membrane constituent is inhibited. Preferred anionic groups are sulfonates and sulfates. Preferred carrier molecules include carbohydrates, polymers, peptides, peptide derivatives, aliphatic groups, alicyclic groups, heterocyclic groups, aromatic groups and combinations thereof.

58 Claims, 10 Drawing figures

Exemplary Claim Number: 1

Number of Drawing Sheets: 8

Full  Title  Citation  Front  Review  Classification  Date  Reference     Claims  KOMC  Drawn Des.

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22. Document ID: US 5869469 A

L19: Entry 22 of 27

File: USPT

Feb 9, 1999

US-PAT-NO: 5869469

DOCUMENT-IDENTIFIER: US 5869469 A

TITLE: Phosphonocarboxylate compounds for treating amyloidosis

DATE-ISSUED: February 9, 1999

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Szarek; Walter A.	Kingston			CA
Kong; Xianqi	Kingston			CA

US-CL-CURRENT: 514/120

## ABSTRACT:

Therapeutic compounds and methods for modulating amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is modulated by the administration to a subject of an effective amount of a therapeutic compound comprising a phosphonate group and a carboxylate group, or a pharmaceutically acceptable salt or ester thereof. In preferred embodiments, an interaction between an amyloidogenic protein and a basement membrane constituent is modulated.

25 Claims, 0 Drawing figures

Exemplary Claim Number: 1

Full  Title  Citation  Front  Review  Classification  Date  Reference     Claims  KOMC  Drawn Des.

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23. Document ID: US 5858326 A

L19: Entry 23 of 27

File: USPT

Jan 12, 1999

h e b b g e e e f e b ef b e

US-PAT-NO: 5858326

DOCUMENT-IDENTIFIER: US 5858326 A

TITLE: Methods of increasing amyloid deposition

DATE-ISSUED: January 12, 1999

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Kisilevsky; Robert	Kingston			CA
Szarek; Walter	Kingston			CA
Weaver; Donald	Kingston			CA
Fraser; Paul	Toronto			CA
Kong; Xianqi	Kingston			CA

US-CL-CURRENT: 424/9.2; 424/78.31, 424/78.35, 435/7.8, 435/7.92, 435/7.93, 435/7.95, 514/772.4, 530/350, 800/9

## ABSTRACT:

In vivo and in vitro methods of increasing amyloid deposition using amyloid-enhancing compounds are described. Methods of forming amyloid fibrils and screening for agents useful in treating amyloidosis are also described. Animals having non-naturally occurring amyloid deposits produced using the amyloid-enhancing compounds even further are described.

5 Claims, 2 Drawing figures

Exemplary Claim Number: 5

Number of Drawing Sheets: 2

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Abstract](#) | [Claims](#) | [KOMC](#) | [Drawn Desc](#)

## □ 24. Document ID: US 5840294 A

L19: Entry 24 of 27

File: USPT

Nov 24, 1998

US-PAT-NO: 5840294

DOCUMENT-IDENTIFIER: US 5840294 A

\*\* See image for Certificate of Correction \*\*

TITLE: Method for treating amyloidosis

DATE-ISSUED: November 24, 1998

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Kisilevsky; Robert	Kingston			CA
Szarek; Walter	Kingston			CA
Weaver; Donald	Kingston			CA

US-CL-CURRENT: 424/78.31; 424/423, 424/427, 424/430, 424/434, 424/436, 424/441, 424/450, 424/78.35, 514/772.4, 526/286, 526/287

## ABSTRACT:

h e b b g e e e f e b ef b e

Therapeutic compounds and methods for inhibiting amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is inhibited by the administration to a subject of an effective amount of a therapeutic compound comprising an anionic group and a carrier molecule, or a pharmaceutically acceptable salt thereof, such that an interaction between an amyloidogenic protein and a basement membrane constituent is inhibited. Preferred anionic groups are sulfonates and sulfates. Preferred carrier molecules include carbohydrates, polymers, peptides, peptide derivatives, aliphatic groups, alicyclic groups, heterocyclic groups, aromatic groups and combinations thereof.

66 Claims, 14 Drawing figures

Exemplary Claim Number: 1

Number of Drawing Sheets: 12

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Claims](#) | [KOMC](#) | [Draw. Desc](#)

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25. Document ID: US 5728375 A

L19: Entry 25 of 27

File: USPT

Mar 17, 1998

US-PAT-NO: 5728375

DOCUMENT-IDENTIFIER: US 5728375 A

TITLE: Method for treating amyloidosis

DATE-ISSUED: March 17, 1998

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Kisilevsky; Robert	Kingston			CA
Szarek; Walter	Kingston			CA
Weaver; Donald	Kingston			CA

US-CL-CURRENT: 424/78.31; 424/450, 424/78.35, 514/772.4, 526/286, 526/287

ABSTRACT:

Therapeutic compounds and methods for inhibiting amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is inhibited by the administration to a subject of an effective amount of a therapeutic compound comprising an anionic group and a carrier molecule, or a pharmaceutically acceptable salt thereof, such that an interaction between an amyloidogenic protein and a basement membrane constituent is inhibited. Preferred anionic groups are sulfonates and sulfates. Preferred carrier molecules include carbohydrates, polymers, peptides, peptide derivatives, aliphatic groups, alicyclic groups, heterocyclic groups, aromatic groups and combinations thereof.

71 Claims, 12 Drawing figures

Exemplary Claim Number: 1

Number of Drawing Sheets: 12

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Claims](#) | [KOMC](#) | [Draw. Desc](#)

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26. Document ID: US 5643562 A

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L19: Entry 26 of 27

File: USPT

Jul 1, 1997

US-PAT-NO: 5643562

DOCUMENT-IDENTIFIER: US 5643562 A

TITLE: Method for treating amyloidosis

DATE-ISSUED: July 1, 1997

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Kisilevsky; Robert	Kingston			CA
Szarek; Walter	Kingston			CA
Weaver; Donald	Kingston			CA

US-CL-CURRENT: 424/78.31; 424/423, 424/427, 424/430, 424/434, 424/436, 424/441,  
424/78.35, 514/772.4, 526/286, 526/287

## ABSTRACT:

Therapeutic compounds and methods for inhibiting amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is inhibited by the administration to a subject of an effective amount of a therapeutic compound comprising an anionic group and a carrier molecule, or a pharmaceutically acceptable salt thereof, such that an interaction between an amyloidogenic protein and a basement membrane constituent is inhibited. Preferred anionic groups are sulfonates and sulfates. Preferred carrier molecules include carbohydrates, polymers, peptides, peptide derivatives, aliphatic groups, alicyclic groups, heterocyclic groups, aromatic groups and combinations thereof.

55 Claims, 12 Drawing figures

Exemplary Claim Number: 1

Number of Drawing Sheets: 12

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KMPC	Drawn Desc
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 27. Document ID: US 5276059 A

L19: Entry 27 of 27

File: USPT

Jan 4, 1994

US-PAT-NO: 5276059

DOCUMENT-IDENTIFIER: US 5276059 A

\*\* See image for Certificate of Correction \*\*

TITLE: Inhibition of diseases associated with amyloid formation

DATE-ISSUED: January 4, 1994

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Caughey; Byron	Hamilton	MT		
Race; Richard	Hamilton	MT		

US-CL-CURRENT: 514/647

h e b b g e e e f

e b ef b e

## ABSTRACT:

The invention provides a method of treating a mammal having a condition associated with formation of amyloidogenic protein without deposition of amyloid plaques. This treatment includes administering to the mammal a pharmacologically effective amount of Congo Red or a pharmaceutically acceptable salt or derivative thereof to interfere with amyloidogenic protein formation or to destabilize amyloidogenic protein structures already formed in said mammal. The invention also provides a method of treating a mammal having a condition associated with deposition of amyloidogenic protein in plaques, and a method of inhibiting the transformation of PrP-sen to PrP-res in a tissue culture sample containing PrP-sen.

34 Claims, 4 Drawing figures

Exemplary Claim Number: 1

Number of Drawing Sheets: 3

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Abstract](#) | [Claims](#) | [KMC](#) | [Drawn Desc](#)

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PMID: 14745700 [PubMed - indexed for MEDLINE]

**2:** [Zanusso G, Casalone C, Acutis P, Bozzetta E, Farinazzo A, Gelati M, Fiorini M, Forloni G, Sv MS, Monaco S, Caramelli M.](#) [Related Articles](#), [Links](#)

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**Synthetic peptide vaccines yield monoclonal antibodies to cellular and pathological prion proteins of ruminants.**

*J Gen Virol.* 1998 Apr;79 ( Pt 4):937-45.

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1: Am J Pathol. 2002 Jul;161(1):13-7. [Related Articles](#), [Links](#)

## Immunization delays the onset of prion disease in mice.

**Sigurdsson EM, Brown DR, Daniels M, Kascsak RJ, Kascsak R, Carp R, Meeker HC, Frangione B, Wisniewski T.**

Department of Psychiatry, New York University School of Medicine, New York, New York 10016, USA.

The outbreak of new variant Creutzfeldt-Jakob disease has raised the specter of a potentially large population being at risk to develop this prionosis. None of the prionoses currently have an effective treatment. Recently, vaccination has been shown to be effective in mouse models of another neurodegenerative condition, namely Alzheimer's disease. Here we report that vaccination with recombinant mouse prion protein delays the onset of prion disease in mice. Vaccination was performed both before peripheral prion exposure and after exposure. A delay in disease onset was seen in both groups, but was more prolonged in animals immunized before exposure. The increase in the incubation period closely correlated with the anti-prion protein antibody titer. This promising finding suggests that a similar approach may work in humans or other mammalian species at risk for prion disease.

PMID: 12107084 [PubMed - indexed for MEDLINE]

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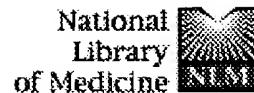
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**7:** [Styrkarsdottir U, Cazier JB, Kong A, Rolfsson O, Larsen H, Bjarnadottir E, Johannsdottir VD, Sigurdardottir MS, Bagger Y, Christiansen C, Reynisdottir I, Grant SF, Jonasson K, Frigge ML, Gulcher JR, Sigurdsson G, Stefansson K.](#) [Related Articles](#), [Links](#)  
**8:** [Linkage of Osteoporosis to Chromosome 20p12 and Association to BMP2.](#) *PLoS Biol.* 2003 Dec;1(3):E69. Epub 2003 Nov 03.  
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**9:** [Stefansson E, Zetterstrom C, Ehlers N, Kiilgaard JF, la Cour M, Sigurdsson H, Gudmundsdottir E, Prause JU, Heijl A.](#) [Related Articles](#), [Links](#)  
**10:** [Nordic research in ophthalmology.](#) *Acta Ophthalmol Scand.* 2003 Dec;81(6):556-66. Review.  
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 Quantitative evaluation by minisequencing and microarrays reveals accurate multiplexed SNP genotyping of whole genome amplified DNA.  
Nucleic Acids Res. 2003 Nov 1;31(21):e129.  
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 **10:** Schiemann O, Fritscher J, Kisseleva N, Sigurdsson ST, Prisner TE. [Related Articles](#), [Links](#)  
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 Copper chelation delays the onset of prion disease.  
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Science. 2003 Jul 11;301(5630):193-6.  
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 Long-term potentiation in freely moving rats reveals asymmetries in thalamic and cortical inputs to the lateral amygdala.  
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Acta Ophthalmol Scand. 2003 Jun;81(3):299-303.  
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23: [Sigurdsson G, Yannopoulos D, McKnite SH, Lurie KG](#). [Related Articles](#), [Links](#)  
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Curr Opin Crit Care. 2003 Jun;9(3):183-8. Review.  
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27: [Gudmundsdottir T, Tryggvadottir L, Allende M, Mast TC, Briem H, Sigurdsson K](#). [Related Articles](#), [Links](#)

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**28: Sigurdsson S, Stromberg R.** Related Articles, Links  
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## Anti-prion antibodies for prophylaxis following prion exposure in mice.

**Sigurdsson EM, Sy MS, Li R, Scholtzova H, Kacsak RJ, Kacsak R, Carp R, Meeker HC, Frangione B, Wisniewski T.**

Department of Psychiatry, New York University School of Medicine, Millhauser Laboratory, 550 First Avenue, New York, NY 10016, USA.

Prion disease is characterized by a conformational change of the normal form of the prion protein (PrP(C)) to the scrapie-associated form (PrP(Sc)). Since the emergence of new variant Creutzfeldt-Jakob disease a potentially large human population is at risk for developing prion disease. Currently, no effective treatment or form of post-exposure prophylaxis is available for prion disease. We recently showed that active immunization with recombinant PrP prolongs the incubation period of scrapie. Here we show that anti-PrP antibodies following prion exposure are effective at increasing the incubation period of the infection. Stimulation of the immune system is an important therapeutic target for the prion diseases, as well as for other neurodegenerative illnesses characterized by abnormal protein conformation.

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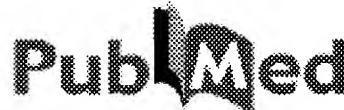
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**Harmeyer S, Pfaff E, Groschup MH.**

Federal Research Centre for Virus Diseases of Animals, Tübingen, Germany.

Transmissible spongiform encephalopathies are closely linked to the accumulation of a pathological isoform of a host-encoded prion protein (PrP (C)), designated PrP(Sc). In an attempt to generate mono- and polyclonal antibodies to ruminant PrP, 32 mice were vaccinated with peptide vaccines which were synthesized according to the amino acid sequence of ovine PrP. By this approach five PrP-reactive polyclonal antisera directed against four different domains of the protein were stimulated. Splenocytes of mice which had developed PrP-reactive antibodies were used for the generation of monoclonal antibodies (MAbs). Obtained PrP-specific MAbs were directed to three different domains of ruminant PrP which differed from the three previously described major MAb binding sites in rodent PrP. MAbs exhibited reactivity with non-denatured ruminant PrP(C) in ELISA and immunoprecipitation and with denatured ovine and bovine PrP(Sc) in immunoblot. Cross-reactivity was observed with PrP(C) of nine other mammalian species and with pathological PrP preferably of ruminants and weakly with that of hamster and mouse. The generated MAbs will be useful tools for the development of diagnostic tests for BSE and scrapie as well as for pathogenesis studies of these diseases.

PMID: 9568991 [PubMed - indexed for MEDLINE]

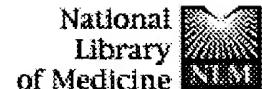
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=> S PrPSc OR ASCR  
34 FILES SEARCHED...  
62 FILES SEARCHED...  
L1 11834 PRPSC OR ASCR

=> S L1 AND adjuvant  
49 FILES SEARCHED...  
L2 108 L1 AND ADJUVANT

=> DUP REM L2  
DUPLICATE IS NOT AVAILABLE IN 'ADISINSIGHT, ADISNEWS, BIOMERCE, DGENE,  
DRUGMONOG2, IMSRESEARCH, FEDRIP, FOREGE, GENBANK, IMSPRODUCT, KOSMET,  
MEDICONF, NUTRACEUT, PCTGEN, PHAR, PHARMAML, PROUSDDR, RDISCLOSURE, SYNTHLINE'.  
ANSWERS FROM THESE FILES WILL BE CONSIDERED UNIQUE  
PROCESSING COMPLETED FOR L2  
L3 76 DUP REM L2 (32 DUPLICATES REMOVED)

=> D L3 1-76

L3 ANSWER 1 OF 76 USPATFULL on STN DUPLICATE 1  
AN 2004:18345 USPATFULL  
TI Methods and compositions for treating a plaque-forming disease  
IN Solomon, Beka, Herzlia Pitauch, ISRAEL  
Frenkel, Dan, Rehovot, ISRAEL  
PA Ramot at Tel-Aviv University Ltd., Tel-Aviv, ISRAEL (non-U.S.  
corporation)  
PI US 2004013647 A1 20040122  
AI US 2003-384788 A1 20030311 (10)  
RLI Continuation-in-part of Ser. No. US 2001-808037, filed on 15 Mar 2001,  
ABANDONED Continuation-in-part of Ser. No. US 2001-830954, filed on 7  
Aug 2001, PENDING Continuation-in-part of Ser. No. US 2002-162889, filed  
on 6 Jun 2002, PENDING Continuation-in-part of Ser. No. US 1999-473653,  
filed on 29 Dec 1999, ABANDONED Continuation-in-part of Ser. No. US  
2000-629971, filed on 31 Jul 2000, ABANDONED Continuation-in-part of  
Ser. No. US 1999-473653, filed on 29 Dec 1999, ABANDONED A 371 of  
International Ser. No. WO 2000-IL518, filed on 31 Aug 2000, PENDING  
Continuation-in-part of Ser. No. US 1999-473653, filed on 29 Dec 1999,  
ABANDONED Continuation-in-part of Ser. No. US 2000-629971, filed on 31  
Jul 2000, ABANDONED Continuation of Ser. No. US 2000-629971, filed on 31  
Jul 2000, ABANDONED  
PRAI US 2002-371735P 20020412 (60)  
US 1999-152417P 19990903 (60)  
US 1999-152417P 19990903 (60)  
DT Utility  
FS APPLICATION  
LN.CNT 4217  
INCL INCLM: 424/093.200  
INCLS: 514/044.000  
NCL NCLM: 424/093.200  
NCLS: 514/044.000  
IC [7]  
ICM: A61K048-00  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 2 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 2004:473109 CAPLUS  
DN 141:37595  
TI M cell directed vaccines  
IN Pascual, David W.  
PA USA  
SO U.S. Pat. Appl. Publ., 33 pp., Cont.-in-part of U.S. Pat. Appl. 2004  
33,486.  
CODEN: USXXCO

DT Patent  
LA English

FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2004109871	A1	20040610	US 2003-660787	20030912
	WO 2001049867	A1	20010712	WO 2001-US426	20010108
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK,				

MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ,  
TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ,

MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,  
DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,  
BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

WO 2002072015 A2 20020919 WO 2002-US7254 20020312

WO 2002072015 A3 20021212

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,  
CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,  
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,  
LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,  
PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,  
UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU,  
TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH,  
CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR,  
BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2004033486 A1 20040219 US 2002-169492 20021021

PRAI US 2000-174786P P 20000106

WO 2001-US426 W 20010108

US 2001-274639P P 20010312

WO 2002-US7254 A2 20020312

US 2002-169492 A2 20021021

L3 ANSWER 3 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN

AN 2004:141667 CAPLUS

DN 140:198062

TI Antigen arrays and compositions comprising antigenic determinant of RANKL  
protein and virus-like particle for treatment of bone disease and prion  
disease

IN Bachmann, Martin; Maurer, Patrick; Spohn, Gunther

PA Switz.

SO U.S. Pat. Appl. Publ., 53 pp., Cont.-in-part of Appl. No. PCT/IB02/00166.  
CODEN: USXXCO

DT Patent

LA English

FAN.CNT 9

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2004033211	A1	20040219	US 2002-289456	20021107
	US 2003175290	A1	20030918	US 2002-50902	20020118
	WO 2002056905	A2	20020725	WO 2002-IB166	20020121
	WO 2002056905	A3	20031009		
		W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM		
		RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG		
PRAI	US 2001-331045P	P	20011107		
	US 2002-50902	A2	20020118		
	WO 2002-IB166	A2	20020121		
	US 2002-396635P	P	20020719		
	US 2001-262379P	P	20010119		
	US 2001-288549P	P	20010504		
	US 2001-326998P	P	20011005		

L3 ANSWER 4 OF 76 USPATFULL on STN

AN 2004:171948 USPATFULL

TI Method

IN Enari, Masato, Chuo-ku, JAPAN  
Flechsig, Eckhard, Versbacher, GERMANY, FEDERAL REPUBLIC OF  
Collinge, John, Queen, UNITED KINGDOM  
Weismann, Charles, London, UNITED KINGDOM

PI US 2004132109 A1 20040708

AI US 2004-470022 A1 20040109 (10)  
WO 2002-GB257 20020122

PRAI GB 2001-1762 20010123

DT Utility

FS APPLICATION

LN.CNT 3141

INCL INCLM: 435/007.200  
INCL INCLS: 435/287.200  
NCL NCLM: 435/007.200  
NCL NCLS: 435/287.200

IC [7]  
ICM: G01N033-53  
ICS: G01N033-567; C12M001-34

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 5 OF 76 USPATFULL on STN

AN 2004:94784 USPATFULL

TI \*\*\*PrPSc\*\*\* -interacting molecules and uses thereof  
IN Cashman, Neil, Toronto, CANADA

Paramithiotis, Eustache, Boucherville, CANADA  
La Boissiere, Sylvie, Montreal, CANADA  
Lawton, Robert, Gorham, ME, UNITED STATES  
Francoeur, Greg, North Yarmouth, ME, UNITED STATES  
Francoeur, Susan, Portland, ME, UNITED STATES LR  
Estey, Lisa, Westbrook, ME, UNITED STATES  
Pinard, Marc, Montreal, CANADA

PI US 2004072236 A1 20040415

AI US 2002-256538 A1 20020927 (10)

DT Utility

FS APPLICATION

LN.CNT 1436

INCL INCLM: 435/007.100

NCL NCLM: 435/007.100

IC [7]  
ICM: G01N033-53

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 6 OF 76 USPATFULL on STN

AN 2004:69593 USPATFULL

TI Fusion proteins comprising DP-178 and other viral fusion inhibitor peptides useful for treating aids

IN Bolognesi, Dani Paul, Durham, NC, UNITED STATES  
Matthews, Thomas James, Durham, NC, UNITED STATES  
Wild, Carl T., Durham, NC, UNITED STATES  
Barney, Shawn O'Lin, Cary, NC, UNITED STATES  
Lambert, Dennis Michael, Cary, NC, UNITED STATES  
Petteway, Stephen Robert, Cary, NC, UNITED STATES  
Langlois, Alphonse J., Durham, NC, UNITED STATES

PA Duke University (U.S. corporation)

Trimeris, Inc. (U.S. corporation)

PI US 2004052820 A1 20040318

AI US 2002-267748 A1 20021008 (10)

RLI Continuation of Ser. No. US 1995-484223, filed on 7 Jun 1995, PENDING  
Division of Ser. No. US 1995-470896, filed on 6 Jun 1995, GRANTED, Pat. No. US 6479055 Continuation-in-part of Ser. No. US 1994-360107, filed on 20 Dec 1994, GRANTED, Pat. No. US 6017536 Continuation-in-part of Ser. No. US 1994-255208, filed on 7 Jun 1994, GRANTED, Pat. No. US 6440656 Continuation-in-part of Ser. No. US 1993-73028, filed on 7 Jun 1993, GRANTED, Pat. No. US 5464933

DT Utility

FS APPLICATION

LN.CNT 40442

INCL INCLM: 424/208.100

INCL INCLS: 424/188.100; 530/350.000; 424/204.100; 530/300.000

NCL NCLM: 424/208.100

NCL NCLS: 424/188.100; 530/350.000; 424/204.100; 530/300.000

IC [7]

ICM: A61K039-21

ICS: C07K014-16; A61K039-12; C07K002-00; C07K004-00; C07K005-00; C07K007-00; C07K014-00; C07K016-00; C07K017-00; A61K038-00; C07K001-00

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 7 OF 76 USPATFULL on STN

AN 2004:69539 USPATFULL

TI Immunization against amyloid plaques using display technology

IN Solomon, Beka, Herzlia Pituach, ISRAEL

Frenkel, Dan, Rehovot, ISRAEL

PA Ramot at Tel-Aviv University, Ltd., Tel Aviv, ISRAEL (non-U.S. corporation)

PI US 2004052766 A1 20040318

AI US 2003-618856 A1 20030715 (10)

RLI Continuation of Ser. No. US 1999-473653, filed on 29 Dec 1999, ABANDONED

PRAI US 1999-152417P 19990903 (60)

DT Utility

FS APPLICATION

LN.CNT 2570

INCL INCLM: 424/093.200

NCL NCLM: 424/093.200

IC [7]

ICM: A61K048-00

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 8 OF 76 USPATFULL on STN

AN 2004:44245 USPATFULL

TI Nucleic acids encoding DP-178 and other viral fusion inhibitor peptides useful for treating aids

IN Bolognesi, Dani Paul, Durham, NC, UNITED STATES

Matthews, Thomas James, Durham, NC, UNITED STATES

Wild, Carl T., Durham, NC, UNITED STATES

PA Duke University (U.S. corporation)

PI US 2004033235 A1 20040219

AI US 2003-267682 A1 20030106 (10)

RLI Continuation of Ser. No. US 1995-484223, filed on 7 Jun 1995, PENDING  
Division of Ser. No. US 1995-470896, filed on 6 Jun 1995, GRANTED, Pat. No. US 6479055 Continuation-in-part of Ser. No. US 1994-360107, filed on 20 Dec 1994, GRANTED, Pat. No. US 6017536 Continuation-in-part of Ser. No. US 1994-255208, filed on 7 Jun 1994, GRANTED, Pat. No. US 6440656 Continuation-in-part of Ser. No. US 1993-73028, filed on 7 Jun 1993, GRANTED, Pat. No. US 5464933

DT Utility

FS APPLICATION

LN.CNT 59510

INCL INCLM: 424/186.100

INCLS: 424/188.100; 530/350.000; 424/208.100; 424/187.100

NCL NCLM: 424/186.100

NCLS: 424/188.100; 530/350.000; 424/208.100; 424/187.100

IC [7]

ICM: A61K039-21

ICS: A61K039-12; C07K014-16; C07K014-10; C07K014-05; C07K014-11

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 9 OF 76 USPATFULL on STN

AN 2004:181084 USPATFULL

TI Immunological detection of prions

IN Korth, Carsten, San Francisco, CA, United States

Stierli, Beat, Daenikon, SWITZERLAND

Stregt, Peter, Zurich, SWITZERLAND

Oesch, Bruno, Stilli, SWITZERLAND

Moser, Markus, Zurich, SWITZERLAND

PA Universitat Zurich, Zurich, SWITZERLAND (non-U.S. corporation)

PI US 6765088 B1 20040720

WO 9837210 19980827

AI US 1999-380015 19990823 (9)

WO 1998-EP917 19980218

PRAI DE 1997-102837 19970221

DT Utility

FS GRANTED

LN.CNT 1703

INCL INCLM: 530/388.100

INCLS: 424/139.100; 424/141.100; 435/007.100; 435/326.000; 435/331.000;  
530/388.850

NCL NCLM: 530/388.100

NCLS: 424/139.100; 424/141.100; 435/007.100; 435/326.000; 435/331.000;  
530/388.850

IC [7]

ICM: C07K016-00

EXF 435/7.1; 435/326; 435/331; 436/503; 436/518; 436/547; 530/387.1;  
530/387.9; 530/388.1; 530/388.85; 800/4; 800/5; 800/6

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 10 OF 76 WPIDS COPYRIGHT 2004 THOMSON DERWENT on STN

AN 2004-501381 [48] WPIDS

DNN N2004-395892 DNC C2004-185869

TI Amplifying immunodetection of pathological prion protein, useful for diagnosis of spongiform encephalopathy, by adding a macrocyclic \*\*\*adjuvant\*\*\* ligand before reaction with antibody.

DC A26 A89 B04 D16 E19 S03

IN COLEMAN, A W; DA SILVA, E; MARTIN, A; MOUSSA, A; SHAHGALDIAN, P; DUPIN, M;

PA LAZAR, A N; LECLERE, E; PERRON, H  
(FRSE-N) AGENCE FR SECURITE SANITAIRES ALIMENTS; (CNRS) CNRS CENT NAT RECH  
SCI; (UYLY-N) UNIV LYON 1 BERNARD CLAUDE; (INMR) BIOMERIEUX SA

CYC 107

PI FR 2849205 A1 20040625 (200448)\* 31 G01N033-68  
WO 2004059322 A1 20040715 (200448) FR G01N033-68  
RW: AT BE BG BW CH CY CZ DE DK EA EE ES FI FR GB GH GM GR HU IE IT KE  
LS LU MC MW MZ NL OA PT RO SD SE SI SK SL SZ TR TZ UG ZM ZW  
W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK  
DM DZ EC EE EG ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP  
KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NI NO NZ OM PG  
PH PL PT RO RU SC SD SE SG SK SL SY TJ TM TN TR TT TZ UA UG US UZ  
VC VN YU ZA ZM ZW

ADT FR 2849205 A1 FR 2002-16383 20021220; WO 2004059322 A1 WO 2003-FR3857  
20031219

PRAI FR 2002-16383 20021220

IC ICM G01N033-68

ICS C12Q001-37; G01N033-566

L3 ANSWER 11 OF 76 DRUGU COPYRIGHT 2004 THOMSON DERWENT on STN  
AN 2004-27090 DRUGU T S

TI Thiotapec-based high-dose chemotherapy with autologous stem-cell rescue in  
patients with recurrent or progressive CNS germ cell tumors.

AU Modak S; Gardner S; Dunkel I J; Balmaceda C; Rosenblum M K; Miller D C;  
Halpern S; Finlay J L

CS Mem.Sloan-Kettering-Cancer-Cent.; Univ.New-York-Columbia; Univ.New-York  
LO New York, N.Y., USA

SO J.Clin.Oncol. (22, No. 10, 1934-43, 2004) 3 Fig. 5 Tab. 40 Ref.  
CODEN: JCONDN ISSN: 0732-183X

AV Department of Pediatrics, Memorial Sloan-Kettering Cancer Center, 1275  
York Ave, New York, NY 10021, U.S.A. (e-mail: modaks@mskcc.org).

LA English

DT Journal

FA AB; LA; CT

FS Literature

L3 ANSWER 12 OF 76 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on  
STN DUPLICATE 2

AN 2004:365298 BIOSIS

DN PREV200400368764

TI Multiple antigenic peptides facilitate generation of anti-prion  
antibodies.

AU Bainbridge, J.; Jones, N.; Walker, B. [Reprint Author]

CS Dept Immunobiol, Natl Inst Biol Stand and Controls, Blanche Lane S Mimms,  
Potters Bar, Herts, EN6 3QG, England  
kbwalker@nibsc.ac.uk

SO Clinical and Experimental Immunology, (August 2004) vol. 137, No. 2, pp.  
298-304. print.

ISSN: 0009-9104 (ISSN print).

DT Article

LA English

ED Entered STN: 8 Sep 2004

Last Updated on STN: 8 Sep 2004

L3 ANSWER 13 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 3

AN 2003:930741 CAPLUS

DN 140:4064

TI vaccine comprising conjugates of prion protein and recombinant virus-like  
particle carrier for treating prion diseases

IN Bachmann, Martin; Maurer, Patrik; Pellicioli, Erica; Renner, Wolfgang A.

PA Cytos Biotechnology Ag, Switz.

SO U.S. Pat. Appl. Publ., 127 pp., Cont.-in-part of Appl. No. PCT/IB02/00166.  
CODEN: USXXCO

DT Patent

LA English

FAN.CNT 9

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI US 2003219459	A1	20031127	US 2003-346190	20030117
US 2003175290	A1	20030918	US 2002-50902	20020118
WO 2002056905	A2	20020725	WO 2002-IB166	20020121
WO 2002056905	A3	20031009		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,  
CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,  
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,  
LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,

PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,  
UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU,

TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH,  
CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR,  
BF, BJ, CF, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

PRAI US 2002-50902 A2 20020118  
WO 2002-IB166 A2 20020121  
US 2002-393725P P 20020708  
US 2002-396590P P 20020718  
US 2001-262379P P 20010119  
US 2001-288549P P 20010504  
US 2001-326998P P 20011005  
US 2001-331045P P 20011107

L3 ANSWER 14 OF 76 IFIPAT COPYRIGHT 2004 IFI on STN DUPLICATE 4

AN 10422134 IFIPAT;IFIUDB;IFICDB

TI SYNTHETIC IMMUNOGENIC BUT NON-DEPOSIT-FORMING POLYPEPTIDES AND PEPTIDES  
HOMOLOGOUS TO AMYLOID BETA, PRION PROTEIN, AMYLIN, ALPHA-SYNUCLEIN, OR  
POLYGLUTAMINE REPEATS FOR INDUCTION OF AN IMMUNE RESPONSE THERETO

IN Frangione Blas; Sigurdsson Einar M; Wisniewski Thomas

PA New York University (59449)

PI US 2003166558 A1 20030904

AI US 2002-301488 20021121

PRAI US 2001-331801P 20011121 (Provisional)

FI US 2003166558 20030904

DT Utility; Patent Application - First Publication

FS CHEMICAL

APPLICATION

CLMN 115

GI 15 Figure(s).

FIG. 1 shows the results of a thioflavin T fluorometric assay. Fibril formation of A beta 1-42, A beta 1-30-NH<sub>2</sub>, and K6A beta 1-30-NH<sub>2</sub> (SEQ ID NO:6) was measured in vitro following incubation at 37 degrees C. K6A beta 1-30-NH<sub>2</sub> was the only peptide that did not form fibrils at any of the time points.

FIGS. 2A and 2B show that A beta 40 and A beta 42 are toxic to human neuroblastoma cells (SK-N-SH) in culture as determined by the MTT assay, whereas K6A beta 30-NH<sub>2</sub> has no effect at 2 days (FIG. 2A) and is slightly trophic at 6 days (FIG. 2B). \*p less than 0.05; \*\*p less than 0.01; \*\*\*p less than 0.001 compared to VEH group (one-way ANOVA).

FIGS. 3A-3D show coronal sections (X50; original magnification) stained with 6E10 against A beta, through the hippocampus and cortex in a Tg control-(FIG. 3A) and K6A beta 1-30-treated (FIG. 3B) Tg mouse. FIGS. 3C and 3D are adjacent sections (X100) double stained for interleukin-1 that recognizes microglia, and A beta. Note the reduction of amyloid burden in the immunized mouse (FIG. 3B), and the lack of ramified microglia (FIG. 3D) surrounding A beta plaque in the same mouse, compared to a control mouse (FIG. 3A, 3C). The bars in FIGS. 3A and 3C are 100 mu m. Abbreviations: hip=hippocampus; cx=cortex; cc=corpus callosum.

FIGS. 4A-4C show the reduction in cortical (FIG. 4A) and hippocampal (FIG. 4B) amyloid burden (6E10) following 7 months treatment with K6A beta 1-30-NH<sub>2</sub>. There is an 89% reduction in cortical amyloid burden (\*p=0.0002; t-test; n=4 per group) and an 81% reduction in hippocampal amyloid burden (\*p=0.0001). Soluble A beta 1-42 levels (FIG. 4C) are reduced by 57% within the brains of the vaccinated mice (\*p=0.0019).

FIG. 5 shows the results of a thioflavin T fluorometric assay. Fibril formation of A beta 1-42, A beta 1-40, A beta 1-30-NH<sub>2</sub>, A beta 1-30K6, A beta 1-30-NH<sub>2</sub> (EE18,19) and A beta 1-30-NH<sub>2</sub> (DD18,19) was measured in vitro following incubation at 37 degrees C. for 15 days. Within this period, no fibril formation of the A beta derivatives containing a polylysine segment or an amino acid substitution within the hydrophobic region was detected.

FIGS. 6A and 6B show the results of MTT cell toxicity assay. Neurotoxicity of A beta 1-42, A beta 1-40, A beta 1-30-NH<sub>2</sub>, K6A beta 1-30-NH<sub>2</sub>, A beta 1-30K6, A beta 1-30-NH<sub>2</sub> (EE,18,19) and A beta 1-30-NH<sub>2</sub> (DD,18,19) was determined following treatment of human neuroblastoma cells (SK-N-SH) for 2 (FIG. 6A) and 6 (FIG. 6B) days. \*p less than 0.05; \*\*p less than 0.01; \*\*\*p less than 0.001 compared to VEH group (one-way ANOVA). In this\*\*\*

\*\*\* assay, A beta 1-40 and A beta 1-42 were toxic to human neuroblastoma\*\*\*

\*\*\* cells (SK-N-SH) in culture. Of the A beta derivatives, even at the\*\*\*

\*\*\* highest concentration (100 mu M), only A beta 1-30K6 displayed a slight\*\*\*

\*\*\* toxicity and only on day 2 of the test. Several of the peptides were\*\*\*

\*\*\* neurotrophic following 6 days incubation. \*p less than 0.05; \*\*p\*\*\*

\*\*\* less than 0.01; \*\*\*p less than 0.001 (One-way Anova; Neuman Keuls' posthoc\*\*\*

\*\*\* test). \*\*\*

\*\*\* FIG. 7 shows the antibody titer determined by ELISA in mice 14 weeks after\*\*\*  
\*\*\* vaccination with mouse recPrP. \*\*\*  
\*\*\* FIGS. 8A and 8B show that a higher anti-PrPC (ME7 FAS PrP) antibody titer\*\*\*  
\*\*\* in vaccinated mice, as presented in FIG. 7, correlates with a longer\*\*\*  
\*\*\* incubation time in both \*\*\*PrPSc\*\*\* inoculated mouse groups at lower  
dilution (FIG. 8A;  $r^2=0.4389$ ,  $p=0.0052$ ) and at higher dilution (FIG. 8B;  
 $r^2=0.6786$ ,  $p$  less than 0.0001).

FIG. 9 is a graph showing the effect of recPrP vaccination on disease onset, with day 0 being the first day an animal scored positive for disease. Group 1 mice were controls inoculated with \*\*\*PrPSc\*\*\* at a 10 fold dilution, while group 2 was inoculated at the same dilution but also received recPrP vaccination. Group 3 mice were controls inoculated with \*\*\*PrPSc\*\*\* at a 1000 fold dilution, while Group 4 received the same dilution of \*\*\*PrPSc\*\*\* along with recPrP vaccination. The two control groups received \*\*\*adjuvant\*\*\* and vehicle injections. Two way ANOVA shows a significant effect for vaccination ( $p=0.0005$ ) and \*\*\*PrPSc\*\*\* dilution ( $p$  less than 0.000001). The Newman-Keuls post-hoc test showed vaccination to have a stronger effect in the 10 fold dilution group (Group 1 versus 2,  $p=0.001$  two-tailed; Group 3 versus 4,  $p=0.036$  one-tailed).

FIG. 10 shows an alignment of amino acid sequences of prion protein (PrP) from human (SEQ ID NO:21), gorilla (SEQ ID NO:22), chimpanzee (SEQ ID NO:23), mouse (SEQ ID NO:24), rat (SEQ ID NO:25), Syrian hamster (SEQ ID NO:26), mink (SEQ ID NO:27), sheep (SEQ ID NO:28), goat (SEQ ID NO:29), cow (SEQ ID NO:30), and greater kudu (SEQ ID NO:31). Amino acid residues that are identical and conserved among the prion proteins of the species presented in this figure are boxed.

FIGS. 11A-C show ELISA evaluation of sera from individual animals vaccinated with K6A beta 1-30-NH2 and alum \*\*\*adjuvant\*\*\*, testing for antibody titer against antigen (FIG. 11A), A beta 142 (FIG. 11B) and A beta 1-40 (FIG. 11C).

FIGS. 12A-C show ELISA evaluation of sera from individual animals immunized with A beta 1-42 and alum \*\*\*adjuvant\*\*\*, testing for antibody titer against antigen (FIG. 12A), K6AP1-30-NH2 (FIG. 12B) and A beta 1-40 (FIG. 12C).

FIGS. 13A and 13B depict a linear maze used to evaluate cognitive capabilities of animals vaccinated with A beta 1-30NH2 and K6A beta 1-30-NH2 together with alum \*\*\*adjuvants\*\*\*, as well as controls.

FIG. 13A shows the maze design during the adaptation phase, and FIG. 13B during testing. Dotted lines indicate blocked alleys.

FIGS. 14A-C depict results obtained from behavioral studies of animals of about 3-4 months of age, after vaccination with A beta 1-30-NH2 and K6A beta 1-30-NH2 together with alum \*\*\*adjuvants\*\*\*, as well as controls. The studies included testing of locomotor activity (FIG. 14A), spontaneous avoidance (FIG. 14B), and passive avoidance (FIG. 14C). See Example 6.

FIGS. 15A-N depict results obtained from behavioral studies of animals of about 11 months of age, after vaccination with A beta 1-30-NH2 and K6A beta 1-30-NH2 together with alum \*\*\*adjuvants\*\*\*, as well as controls. The studies included testing of locomotor activity (FIG. 15A), and cognitive testing using traverse beam (FIGS. 15B and 15C), rotarod (FIG. 15D), radial arm maze (FIGS. 15E and 15F), straight alley channel (FIG. 15G), visible platform (FIGS. 15H and 15T), Morris water maze (FIGS. 15J and 15K), probe trial (FIGS. 15L and 15M), and linear maze (FIG. 15N). See Example 6.

L3 ANSWER 15 OF 76 IFIPAT COPYRIGHT 2004 IFI on STN DUPLICATE 5  
AN 10304838 IFIPAT;IFIUDB;IFICDB  
TI AGENT; THERAPY, PREVENTION PRION INFECTIONS; USING MONOCLONAL ANTIBODIES  
ENCAPSULATED AS HYBRIDOMAS  
IN Enari Masato (JP); Weissmann Charles (GB)  
PA Unassigned Or Assigned To Individual (68000)  
PI US 2003049249 A1 20030313  
AI US 2001-985164 20011101  
PRAI GB 2001-221621 20010913  
FI US 2003049249 20030313  
DT Utility; Patent Application - First Publication  
FS CHEMICAL  
APPLICATION  
OS CA 138:231762  
CLMN 8  
GI 4 Figure(s).

FIG. 1 represents the susceptibility to scrapie infection and PrP level of various sublines of N2a cells. N2a populations as propagated routinely in the lab and single clones transformed with a PrP expression or a control vector are seeded into 24well plates (2 x 104 cells/well) and grown to

confluence. (a) Cultures are exposed for 3 days to purified mouse \*\*\*PrPSc\*\*\* (RML strain, 20 ng/ml), cultured for 29 days (8 passages) and assayed for \*\*\*PrPSc\*\*\* formation by the cell blot assay. (b) Prionsusceptible N2a/Bos2 and resistant N2a/2M11 cells are exposed for 3 days to the dilutions indicated of infected 10% brain homogenate, cultured for 14 days (3 passages) and assayed for \*\*\*PrPSc\*\*\* formation. Cells exposed to a 10<sup>-4</sup> dilution are still slightly positive (c) Western blot analysis of N2a sublines is performed using monoclonal anti-PrP antibody 6H4. Cells transfected with the expression plasmid for mouse PrPc, MHM2 PrP or MH2M PrP are indicated by mo, M2 or 2M, respectively. BOS designates cells cotransfected with pSVneo and pEF-BOS-EX. N2a, the original uncloned cells, as well as the highly susceptible N2a/Bos2 cell line show similar, low expression of PrPc as compared to the non-susceptible moS or 2M11 lines.

FIG. 2 represents anti PrP antibody 6H4 and PIPLC preventing infection of N2a/Bos2 cell with scrapie prions and abrogate \*\*\*PrPSc\*\*\* accumulation in chronically infected cells. (a) N2a/Bos2 cells are incubated for 2 h with antibody 6H4 or PIPLC at the concentrations indicated and exposed to 0.1% scrapie-infected brain homogenate (fmal concentration) for 3 days. After culturing for 14 days (3 passages) in the absence of PIPLC or in the continued presence of 6H4, \*\*\*PrPSc\*\*\* expression is determined. (b,c) Chronically scrapie-infected N2a/Bos2 cells are cultured for (b) 3 days at the levels of antibody 6H4 or of PIPLC indicated or (c) 14 days (3 passages) at the concentrations of antibody 6H4 indicated, and \*\*\*PrPSc\*\*\* accumulation is monitored. (d) Chronically scrapie-infected N2a/ Bos2 cells are exposed to 6H4 at the concentrations indicated for 2 weeks and further cultured in the absence of the antibody for 6 weeks. Cultures are split 1:5 every 34 days. There is no reappearance of \*\*\*PrPSc\*\*\*. "Cell staining" refers to staining of the membranes with ethidium bromide to monitor efficiency of transfer of the cell layer. IN, chronically scrapie-infected N2a/Bos2 cells.

FIG. 3 represents chronically infected N2a/Bos2 cells "cured" of \*\*\*PrPSc\*\*\* by antibody 6H4 treatment continuing to produce PrP and their susceptibility to reinfection. (a) Chronically infected N2a/Bos2 cells, treated with antibody 6H4 for 2 weeks at the concentrations indicated and propagated in its absence for 66 days were not exposed (top 2 rows) or exposed to 0.1% RMLinfected mouse brain homogenate for 3 days (bottom 2 rows). After culturing for 14 days, \*\*\*PrPSc\*\*\* is monitored by the cell blot assay. (b) Relative susceptibility to prions of N2a/Bos2 cells (BOS2) and N2a/Bos2 cells "cured" by exposure to antibody 6H4 at 20  $\mu$ g/ml is determined by exposing cultures to various dilutions of RML-infected mouse brain homogenate for 3 days and determining \*\*\*PrPSc\*\*\* as above. (c) Levels of PrPc and \*\*\*PrPSc\*\*\* in various sublines are determined by Western blotting. Chronically infected N2a/Bos2 cells, treated for 2 weeks with antibody 6H4 at the concentrations indicated are passaged for 84 days after antibody withdrawal. Cells are lysed and samples corresponding to 2.25 x 10<sup>5</sup> cells are incubated in the presence (+PK) or absence (-PK) of proteinase K (5  $\mu$ g/ml) for 90 minutes at 37 degrees C. Western blotting is performed as described in Methods. UN, uninfected N2a/Bos2 cells and I-BOS2, chronically prion-infected BOS2 cells. Molecular weight markers are indicated at the left of each panel.

FIG. 4 represents a model to explain abolition of \*\*\*PrPSc\*\*\* by anti PrP antibody (or PIPLC). PrPc is attached to the membrane by a glycosylphosphatidyl inositol anchor and cycles between the cell surface and an endocytic compartment (43). In scrapieinfected cells, PrPc is recruited into \*\*\*PrPSc\*\*\* "seeds" (44), which may be located at the cell surface and/or in the endocytic/lysosomal compartment. \*\*\*PrPSc\*\*\* is degraded with a halflife of about 15 h (37); if PrPc is prevented from converting to \*\*\*PrPSc\*\*\* by either a blocking antibody or by being stripped from the cell surface by PIPLC, \*\*\*PrPSc\*\*\* will diminish and ultimately disappear.

L3 ANSWER 16 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 2004:2727 CAPLUS  
DN 140:75940

TI Unmethylated CpG oligonucleotide-packaged virus-like particles for  
enhancing immune response of vaccines  
IN Bachman, Martin F.; Renner, Wolfgang A.  
PA Cytos Biotechnology Ag, Switz.  
SO PCT Int. Appl., 252 pp.  
CODEN: PIXXD2  
DT Patent  
LA English  
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2004000351	A1	20031231	WO 2003-EP6541	20030620
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
US 2004005338	A1	20040108	US 2003-465811	20030620
PRAI US 2002-389898P	P	20020620		
RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT				

L3	ANSWER 17 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN			
AN	2003:590953 CAPLUS			
DN	139:128014			
TI	Methods for treating diseases or conditions with peptide constructs			
IN	Zimmerman, Daniel H.; Charoenvit, Yupin; Rosenthal, Kenneth S.; Whelan, Mike			
PA	Cel-Sci Corporation, USA			
SO	PCT Int. Appl., 73 pp.			
CODEN: PIXXD2				
DT	Patent			
LA	English			
FAN.CNT 1				
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2003061589	A2	20030731	WO 2003-US1816	20030123
WO 2003061589	A3	20040527		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
PRAI US 2002-349982P	P	20020123		
US 2002-349983P	P	20020123		
US 2002-350032P	P	20020123		

L3	ANSWER 18 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN			
AN	2003:570846 CAPLUS			
DN	139:132445			
TI	Pharmacine comprising conjugates of prion protein and virus-like particle carrier for treating prion diseases			
IN	Bachmann, Martin; Maurer, Patrick; Pellicioli, Erica; Renner, Wolfgang A.			
PA	Cytos Biotechnology A.-G., Switz.			
SO	PCT Int. Appl., 247 pp.			
CODEN: PIXXD2				
DT	Patent			
LA	English			
FAN.CNT 9				

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2003059386	A2	20030724	WO 2003-EP460	20030117
WO 2003059386	A3	20040311		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW,				

ML, MR, NE, SN, TD, TG				
US 2003175290	A1	20030918	US 2002-50902	20020118
WO 2002056905	A2	20020725	WO 2002-IB166	20020121
WO 2002056905	A3	20031009		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
PRAI US 2002-50902	A	20020118		
WO 2002-IB166	A	20020121		
US 2002-393725P	P	20020708		
US 2002-396590P	P	20020718		
US 2001-262379P	P	20010119		
US 2001-288549P	P	20010504		
US 2001-326998P	P	20011005		
US 2001-331045P	P	20011107		

L3 ANSWER 19 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN

AN 2003:434266 CAPLUS

DN 139:21013

TI Synthetic immunogenic/non-deposit-forming polypeptides and peptides homologous to amyloid .beta., prion protein, amylin, .alpha.-synuclein, or polyglutamine repeats for induction of an immune response

IN Frangione, Blas; Wisniewski, Thomas; Sigurdsson, Einar M.

PA New York University, USA

SO PCT Int. Appl., 265 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	-----	-----	-----	-----
PI WO 2003045128	A2	20030605	WO 2002-US37634	20021121
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
US 2003166558	A1	20030904	US 2002-301488	20021121
PRAI US 2001-331801P	P	20011121		

L3 ANSWER 20 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN

AN 2003:334937 CAPLUS

DN 138:349696

TI Fusion protein comprising cytokines, chemokines, and interferons for use as vaccine \*\*\*adjuvant\*\*\* in immunotherapy for cancer and viral infection

IN Galipeau, Jacques; Stagg, John

PA Centre for Translational Research In Cancer, Can.

SO PCT Int. Appl., 53 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	-----	-----	-----	-----
PI WO 2003035105	A2	20030501	WO 2002-CA1649	20021023
WO 2003035105	A3	20030918		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD,				

RU, TJ, TM  
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

EP 1440090 A2 20040728 EP 2002-769821 20021023  
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK  
PRAI US 2001-330476P P 20011023  
WO 2002-CA1649 W 20021023

L3 ANSWER 21 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 2003:242184 CAPLUS  
DN 138:285995  
TI Packaging of immunostimulatory substances and antigens into virus-like particles for use as vaccines against cancer, autoimmune disease, allergy and viral infection

IN Maurer, Patrick; Tissot, Alain; Schwarz, Katrin; Meijerink, Edwin; Lipowsky, Gerad; Pompens, Paul; Cielens, Indulis; Renhofa, Regina; Bachmann, Martin F.; Storni, Tazio  
PA Cytos Biotechnology A.-G., Switz.  
SO PCT Int. Appl., 322 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003024481	A2	20030327	WO 2002-IB4132	20020916
	WO 2003024481	A3	20040603		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	US 2003099668	A1	20030529	US 2002-244065	20020916
	EP 1450856	A2	20040901	EP 2002-777600	20020916
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK				
PRAI	US 2001-318994P	P	20010914		
	US 2002-374145P	P	20020422		
	WO 2002-IB4132	W	20020916		

L3 ANSWER 22 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 2003:242183 CAPLUS  
DN 138:270293  
TI Vaccine compositions comprising anti-CD4 antibody or immunostimulatory nucleic acid and antigen-coupled virus-like particles for enhancement of immune responses

IN Bachmann, Martin F.; Storni, Tazio; Lechner, Franziska  
PA Cytos Biotechnology A.-G., Switz.  
SO PCT Int. Appl., 243 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003024480	A2	20030327	WO 2002-IB4252	20020916
	WO 2003024480	A3	20031030		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR,				

NE, SN, TD, TG  
US 2003091593 A1 20030515 US 2002-243739 20020916  
EP 1425040 A2 20040609 EP 2002-783338 20020916  
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,  
IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK  
PRAI US 2001-318967P P 20010914  
WO 2002-IB4252 W 20020916

L3 ANSWER 23 OF 76 IFIPAT COPYRIGHT 2004 IFI on STN  
AN 10454587 IFIPAT;IFIUDB;IFICDB  
TI FRAGMENTS OF PRION PROTEINS  
IN Fishleigh Robert Vincent (GB); Mee Roger Paul (GB); Robson Barry (GB)  
PA Proteus Molecular Design Ltd GB (43521)  
PI US 2003199013 A1 20031023  
AI US 2002-116061 20020405  
RLI WO 1992-GB2246 19921203 Section 371 PCT Filing UNKNOWN  
US 1994-244701 19940602 DIVISION 5773572  
US 1998-76721 19980513 DIVISION 6379905  
PRAI GB 1991-257477 19911203  
GB 1992-146638 19920710  
FI US 2003199013 20031023  
US 5773572  
US 6379905  
DT Utility; Patent Application - First Publication  
FS CHEMICAL  
APPLICATION  
CLMN 45

L3 ANSWER 24 OF 76 USPATFULL on STN  
AN 2003:334947 USPATFULL  
TI Nucleotide sequences that code for torsin genes, torsin proteins, and  
methods of using the same to treat protein-aggregation  
IN Caldwell, Guy A., Tuscaloosa, AL, UNITED STATES  
Caldwell, Kim A., Tuscaloosa, AL, UNITED STATES  
PA THE UNIVERSITY OF ALABAMA, Tuscaloosa, AL (U.S. corporation)  
PI US 2003235823 A1 20031225  
AI US 2002-177104 A1 20020624 (10)  
DT Utility  
FS APPLICATION  
LN.CNT 3296  
INCL INCLM: 435/006.000  
INCLS: 435/007.100; 435/069.100; 435/320.100; 435/325.000; 530/350.000;  
530/388.100; 536/023.500; 514/012.000  
NCL NCLM: 435/006.000  
NCLS: 435/007.100; 435/069.100; 435/320.100; 435/325.000; 530/350.000;  
530/388.100; 536/023.500; 514/012.000  
IC [7]  
ICM: C12Q001-68  
ICS: G01N033-53; C12P021-02; C12N005-06; C07K014-47; C07K016-20;  
A61K038-17  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 25 OF 76 USPATFULL on STN  
AN 2003:330543 USPATFULL  
TI Immunological methods and compositions for the treatment of Alzheimer's  
disease  
IN St. George-Hyslop, Peter H., Toronto, CANADA  
McLaurin, JoAnne, Toronto, CANADA  
PA Hospital for Sick Children and University of Toronto (non-U.S.  
corporation)  
PI US 2003232758 A1 20031218  
AI US 2003-411544 A1 20030410 (10)  
PRAI US 2002-373914P 20020419 (60)  
DT Utility  
FS APPLICATION  
LN.CNT 2487  
INCL INCLM: 514/012.000  
INCLS: 530/324.000; 435/069.100; 435/320.100; 435/325.000; 536/023.100  
NCL NCLM: 514/012.000  
NCLS: 530/324.000; 435/069.100; 435/320.100; 435/325.000; 536/023.100  
IC [7]  
ICM: A61K038-17  
ICS: C07K014-47; C12P021-02; C12N005-06  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 26 OF 76 USPATFULL on STN

AN 2003:324329 USPATFULL  
TI Antibodies specific for native \*\*\*PrPsc\*\*\*  
IN Prusiner, Stanley B., San Francisco, CA, UNITED STATES  
Williamson, R. Anthony, San Diego, CA, UNITED STATES  
Burton, Dennis R., LaJolla, CA, UNITED STATES  
PA The Regents of the University of California and The Scripps Research  
Institute (U.S. corporation)  
PI US 2003228303 A1 20031211  
AI US 2003-435602 A1 20030509 (10)  
RLI Continuation of Ser. No. US 2001-943906, filed on 30 Aug 2001, GRANTED,  
Pat. No. US 6562341 Continuation of Ser. No. US 2000-550374, filed on 13  
Apr 2000, GRANTED, Pat. No. US 6372214 Continuation of Ser. No. US  
1998-36579, filed on 6 Mar 1998, GRANTED, Pat. No. US 6290954 Division  
of Ser. No. US 1996-713939, filed on 13 Sep 1996, GRANTED, Pat. No. US  
5846533 Continuation-in-part of Ser. No. US 1995-528104, filed on 14 Sep  
1995, ABANDONED

DT Utility

FS APPLICATION

LN.CNT 2983

INCL INCLM: 424/130.100

NCL NCLM: 424/130.100

IC [7]

ICM: A61K039-395

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 27 OF 76 USPATFULL on STN

AN 2003:264865 USPATFULL

TI Therapy for human cancers using cisplatin and other drugs or genes  
encapsulated into liposomes

IN Boulikas, Teni, Palo Alto, CA, UNITED STATES

PI US 2003185879 A1 20031002

AI US 2003-350470 A1 20030123 (10)

RLI Division of Ser. No. US 1999-434345, filed on 5 Nov 1999, GRANTED, Pat.  
No. US 6511676

DT Utility

FS APPLICATION

LN.CNT 1652

INCL INCLM: 424/450.000

INCLS: 424/649.000

NCL NCLM: 424/450.000

NCLS: 424/649.000

IC [7]

ICM: A61K009-127

ICS: A61K033-24

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 28 OF 76 USPATFULL on STN

AN 2003:264794 USPATFULL

TI Prostate cell lines

IN Thraves, Peter, London, UNITED KINGDOM

Sutton, Andrew, London, UNITED KINGDOM

PI US 2003185808 A1 20031002

AI US 2002-240523 A1 20021023 (10)

WO 2001-GB1437 20010330

PRAI GB 2000-8032 20000401

GB 2000-24237 20001003

DT Utility

FS APPLICATION

LN.CNT 727

INCL INCLM: 424/093.210

INCLS: 514/044.000; 435/366.000; 424/085.200

NCL NCLM: 424/093.210

NCLS: 514/044.000; 435/366.000; 424/085.200

IC [7]

ICM: A61K048-00

ICS: C12N005-08

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 29 OF 76 USPATFULL on STN

AN 2003:206867 USPATFULL

TI Antibodies specific for ungulate PrP

IN Prusiner, Stanley B., San Francisco, CA, UNITED STATES

Safar, Jiri G., Walnut Creek, CA, UNITED STATES

Williamson, R. Anthony, San Diego, CA, UNITED STATES

Burton, Dennis R., La Jolla, CA, UNITED STATES

PI US 2003143224 A1 20030731

AI US 2003-355780 A1 20030130 (10)  
RLI Continuation of Ser. No. US 2000-627218, filed on 27 Jul 2000, GRANTED,  
Pat. No. US 6537548  
DT Utility  
FS APPLICATION  
LN.CNT 2123  
INCL INCLM: 424/130.100  
INCLS: 435/345.000; 435/006.000  
NCL NCLM: 424/130.100  
NCLS: 435/345.000; 435/006.000  
IC [7]  
ICM: C12Q001-68  
ICS: A61K039-395; C12N005-06; C12N005-16  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 30 OF 76 USPATFULL on STN  
AN 2003:181684 USPATFULL  
TI Surface simulation synthetic peptides useful in the treatment of  
hyper-variable viral pathogens  
IN Crevecoeur, Harry F., Valley Stream, NY, UNITED STATES  
PI US 2003125518 A1 20030703  
AI US 2001-12806 A1 20011201 (10)  
DT Utility  
FS APPLICATION  
LN.CNT 1650  
INCL INCLM: 530/327.000  
INCLS: 530/328.000; 435/005.000; 702/019.000  
NCL NCLM: 530/327.000  
NCLS: 530/328.000; 435/005.000; 702/019.000  
IC [7]  
ICM: C07K007-08  
ICS: C12Q001-70; G01N033-48; G06F019-00  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 31 OF 76 USPATFULL on STN  
AN 2003:126727 USPATFULL  
TI Novel methods for down-regulation of amyloid  
IN Jensen, Martin Roland, Horsholm, DENMARK  
Birk, Peter, Horsholm, DENMARK  
Nielsen, Klaus Gregorius, Horsholm, DENMARK  
PI US 2003086938 A1 20030508  
AI US 2002-204362 A1 20020816 (10)  
WO 2001-DK113 20010219  
PRAI DK 2000-265 20000221  
DT Utility  
FS APPLICATION  
LN.CNT 3114  
INCL INCLM: 424/185.100  
NCL NCLM: 424/185.100  
IC [7]  
ICM: A61K039-00  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 32 OF 76 USPATFULL on STN  
AN 2003:112522 USPATFULL  
TI Agents and compositions and methods utilizing same useful in diagnosing  
and/or treating or preventing plaque forming  
IN Solomon, Beka, Herzlia Pituach, ISRAEL  
Hanan, Eilat, Tel Aviv, ISRAEL  
Frenkel, Dan, Rehovot, ISRAEL  
PA Ramot University Authority for Applied Research & Industrial  
Development, Tel Aviv, ISRAEL (non-U.S. corporation)  
PI US 2003077252 A1 20030424  
AI US 2002-162889 A1 20020606 (10)  
RLI Continuation of Ser. No. US 2000-629971, filed on 31 Jul 2000, ABANDONED  
Continuation-in-part of Ser. No. US 1999-473653, filed on 29 Dec 1999,  
PENDING  
PRAI US 1999-152417P 19990903 (60)  
DT Utility  
FS APPLICATION  
LN.CNT 2994  
INCL INCLM: 424/093.200  
INCLS: 514/044.000; 435/456.000; 435/235.100  
NCL NCLM: 424/093.200  
NCLS: 514/044.000; 435/456.000; 435/235.100  
IC [7]

ICM: A61K048-00

ICS: C12N007-01; C12N015-86

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 33 OF 76 USPATFULL on STN  
AN 2003:44722 USPATFULL  
TI Early pre-symptomatic prion diagnostic blood test for encephalopathies  
IN Resink, Annelies, Paris, FRANCE  
Fuentes, Nathalie, Kremlin Bicetre, FRANCE  
Schweighoffer, Fabien, Vincennes, FRANCE  
PI US 2003032032 A1 20030213  
AI US 2002-100178 A1 20020319 (10)  
PRAI US 2001-278670P 20010321 (60)  
US 2001-282463P 20010410 (60)  
DT Utility  
FS APPLICATION  
LN.CNT 962  
INCL INCLM: 435/006.000  
NCL NCLM: 435/006.000  
IC [7]  
ICM: C12Q001-68

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 34 OF 76 USPATFULL on STN  
AN 2003:273427 USPATFULL  
TI Inhibitors of amyloid formation  
IN Caughey, Winslow S., Hamilton, MT, United States  
Caughey, Byron, Hamilton, MT, United States  
PA The United States of America as represented by the Department of Health  
and Human Services, Washington, DC, United States (U.S. government)  
PI US 6632808 B1 20031014  
WO 2000009111 20000224  
AI US 2001-762725 20010307 (9)  
WO 1999-US18297 19990811  
PRAI US 1998-96148P 19980811 (60)  
DT Utility  
FS GRANTED  
LN.CNT 1503  
INCL INCLM: 514/185.000  
INCLS: 514/410.000; 540/122.000; 540/145.000  
NCL NCLM: 514/185.000  
NCLS: 514/410.000; 540/122.000; 540/145.000  
IC [7]  
ICM: A61K031-409  
ICS: C07D487-22  
EXF 514/185; 514/410; 540/122; 540/145  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 35 OF 76 USPATFULL on STN  
AN 2003:81453 USPATFULL  
TI Antibodies specific for ungulate PrP  
IN Prusiner, Stanley B., San Francisco, CA, United States  
Safar, Jiri, Concord, CA, United States  
Williamson, R. Anthony, San Diego, CA, United States  
Burton, Dennis R., La Jolla, CA, United States  
PA The Regents of the University of California, Oakland, CA, United States  
(U.S. corporation)  
The Scripps Research Institute, La Jolla, CA, United States (U.S.  
corporation)  
PI US 6537548 B1 20030325  
AI US 2000-627218 20000727 (9)  
DT Utility  
FS GRANTED  
LN.CNT 2073  
INCL INCLM: 424/130.100  
INCLS: 424/009.100; 424/185.100; 435/007.100; 435/070.100; 435/071.100;  
530/387.100; 530/398.100  
NCL NCLM: 424/130.100  
NCLS: 424/009.100; 424/185.100; 435/007.100; 435/070.100; 435/071.100;  
530/387.100; 530/389.100  
IC [7]  
ICM: A61K039-395  
EXF 424/9.1; 424/130.1; 424/185.1; 435/7.1; 435/70.1; 435/71.1; 436/503;  
436/547; 530/387.1; 530/388.27; 530/398.1  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 36 OF 76 USPATFULL on STN  
AN 2003:40533 USPATFULL  
TI Methods for the inhibition of epstein-barr virus transmission employing anti-viral peptides capable of abrogating viral fusion and transmission  
IN Barney, Shawn O'Lin, Cary, NC, United States  
Lambert, Dennis Michael, Cary, NC, United States  
Petteway, Stephen Robert, Cary, NC, United States  
PA Trimeris, Inc., Durham, NC, United States (U.S. corporation)  
PI US 6518013 B1 20030211  
AI US 1995-485546 19950607 (8)  
RLI Continuation-in-part of Ser. No. US 1994-360107, filed on 20 Dec 1994, now patented, Pat. No. US 6017536 Continuation-in-part of Ser. No. US 1994-255208, filed on 7 Jun 1994 Continuation-in-part of Ser. No. US 1993-73028, filed on 7 Jun 1993, now patented, Pat. No. US 5464933  
DT Utility  
FS GRANTED  
LN.CNT 24700  
INCL INCLM: 435/005.000  
INCLS: 424/230.100; 530/300.000; 530/324.000; 530/325.000; 530/326.000  
NCL NCLM: 435/005.000  
NCLS: 424/230.100; 530/300.000; 530/324.000; 530/325.000; 530/326.000  
IC [7]  
ICM: C12Q001-70  
EXF 435/5; 530/300; 530/324-329; 530/350; 424/230.1  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 37 OF 76 USPATFULL on STN  
AN 2003:26157 USPATFULL  
TI Therapy for human cancers using cisplatin and other drugs or genes encapsulated into liposomes  
IN Boulikas, Teni, 249 Matadero Ave., Palo Alto, CA, United States 94306  
PI US 6511676 B1 20030128  
AI US 1999-434345 19991105 (9)  
DT Utility  
FS GRANTED  
LN.CNT 1642  
INCL INCLM: 424/450.000  
INCLS: 264/004.100; 264/004.300  
NCL NCLM: 424/450.000  
NCLS: 264/004.100; 264/004.300  
IC [7]  
ICM: A61K009-127  
EXF 424/450; 264/4.1; 264/4.3  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 38 OF 76 DRUGU COPYRIGHT 2004 THOMSON DERWENT on STN  
AN 2004-07769 DRUGU P  
TI Immunotherapy as a therapeutic treatment for neurodegenerative disorders.  
AU White A R; Hawke S H  
CS Howard-Florey-Inst.; Univ.London  
LO Melbourne, Austr.; London, U.K.  
SO J.Neurochem. (87, No. 4, 801-08, 2003) 1 Fig. 56 Ref.  
CODEN: JONRA9 ISSN: 0022-3042  
AV Neurochemistry Group, Howard Florey Institute of Experimental Physiology and Medicine, Victoria 3010, Australia. (e-mail: a.white@hfi.unimelb.edu.au).  
LA English  
DT Journal  
FA AB; LA; CT  
FS Literature

L3 ANSWER 39 OF 76 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on STN DUPLICATE 6  
AN 2003:86974 BIOSIS  
DN PREV200300086974  
TI Complete Freund's \*\*\*adjuvant\*\*\* immunization prolongs survival in experimental prion disease in mice.  
AU Tal, Yuval; Souan, Lina; Cohen, Irun R. [Reprint Author]; Meiner, Zeev; Taraboulos, Albert; Mor, Felix  
CS Department of Immunology, Weizmann Institute of Science, P.O. Box 26, Rehovot, 76100, Israel  
SO irun.cohen@weizmann.ac.il  
Journal of Neuroscience Research, (January 15 2003) Vol. 71, No. 2, pp. 286-290. print.  
ISSN: 0360-4012 (ISSN print).  
DT Article

LA English  
ED Entered STN: 6 Feb 2003  
Last Updated on STN: 6 Feb 2003

L3 ANSWER 40 OF 76 MEDLINE on STN  
AN 2003475318 MEDLINE  
DN PubMed ID: 14550926  
TI Immunisation with a synthetic prion protein-derived peptide prolongs survival times of mice orally exposed to the scrapie agent.  
AU Schwarz Anja; Kratke Oliver; Burwinkel Michael; Riemer Constanze; Schultz Julia; Henklein Peter; Bamme Theresa; Baier Michael  
CS Project Neurodegenerative Diseases, Robert-Koch-Institute, Nordufer 20, 13353 Berlin, Germany.  
SO Neuroscience letters, (2003 Oct 30) 350 (3) 187-9.  
CY Ireland  
DT Journal; Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals  
EM 200311  
ED Entered STN: 20031011  
Last Updated on STN: 20031111  
Entered Medline: 20031110

L3 ANSWER 41 OF 76 USPATFULL on STN DUPLICATE 7  
AN 2002:272456 USPATFULL  
TI Antibodies specific for native \*\*\*PrPSC\*\*\*  
IN Prusiner, Stanley B., San Francisco, CA, UNITED STATES  
Williamson, R. Anthony, San Diego, CA, UNITED STATES  
Burton, Dennis R., La Jolla, CA, UNITED STATES  
US 2002150571 A1 20021017  
US 6562341 B2 20030513  
US 2001-943906 A1 20010830 (9)  
PI Continuation of Ser. No. US 2000-550374, filed on 13 Apr 2000, PENDING  
AI Continuation of Ser. No. US 1998-36579, filed on 6 Mar 1998, PATENTED  
RLI Division of Ser. No. US 1996-713939, filed on 13 Sep 1996, PATENTED  
Continuation-in-part of Ser. No. US 1995-528104, filed on 14 Sep 1995, ABANDONED  
OT Utility  
FS APPLICATION  
LN.CNT 2374  
INCL INCLM: 424/130.100  
NCL NCLM: 424/130.100  
NCLS: 424/009.100; 424/009.200; 424/147.100; 435/007.100; 435/070.100; 435/071.100; 436/503.000; 436/518.000; 436/547.000; 530/387.100  
IC [7]  
ICM: A61K039-395  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 42 OF 76 USPATFULL on STN DUPLICATE 8  
AN 2002:259529 USPATFULL  
TI Discordant helix stabilization for prevention of amyloid formation  
IN Johansson, Jan, Stockholm, SWEDEN  
PI US 2002143105 A1 20021003  
US 6716589 B2 20040406  
AI US 2001-988842 A1 20011119 (9)  
PRAI US 2000-253695P 20001120 (60)  
US 2000-251662P 20001206 (60)  
OT Utility  
FS APPLICATION  
LN.CNT 1541  
INCL INCLM: 525/054.100  
NCL NCLM: 435/007.200  
IC [7]  
ICM: C08H001-00  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 43 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 2002:185399 CAPLUS  
DN 136:229029  
TI Method for precipitating mono and multiple layers of organophosphoric and organophosphonic acids and the salts thereof in addition to use thereof  
N Hofer, Rolf; Pawlak, Michael; Textor, Marcus; Schuermann-Mader, Eveline; Ehrat, Markus; Tosatti, Samuele  
PA Zepotosens A.-G., Switz.  
O PCT Int. Appl., 88 pp.

CODEN: PIXXD2

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002020873	A2	20020314	WO 2001-EP10077	20010831
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	AU 2001089859	A5	20020322	AU 2001-89859	20010831
	EP 1315968	A2	20030604	EP 2001-969680	20010831
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
PRAI	US 2003186914	A1	20031002	US 2003-363555	20030305
	CH 2000-1732	A	20000905		
	WO 2001-EP10077	W	20010831		
OS	MARPAT 136:229029				

L3 ANSWER 44 OF 76 USPATFULL on STN

AN 2002:329478 USPATFULL

TI Novel method for down-regulation of amyloid

IN Jensen, Martin Roland, Holte, DENMARK

Rasmussen, Peter Birk, Frederiksberg, DENMARK

Nielsen, Klaus Gregorius, Soborg, DENMARK

PI US 2002187157 A1 20021212

AI US 2001-785215 A1 20010220 (9)

PRAI PA 2000-200000265 20000221

US 2000-186295P 20000301 (60)

DT Utility

FS APPLICATION

LN.CNT 3272

INCL INCLM: 424/185.100

INCLS: 424/085.100; 424/085.200

NCL NCLM: 424/185.100

NCLS: 424/085.100; 424/085.200

IC [7]

ICM: A61K039-00

ICS: A61K038-19; A61K038-20

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 45 OF 76 USPATFULL on STN

AN 2002:251957 USPATFULL

TI Conformational and topological protein regulation

IN Lingappa, Vishwanath R., San Francisco, CA, UNITED STATES

Rutkowski, D. Thomas, San Francisco, CO, UNITED STATES

Hegde, Ramanugan S., Rockville, MD, UNITED STATES

PI US 2002137915 A1 20020926

AI US 2000-739179 A1 20001215 (9)

PRAI US 1999-171012P 19991215 (60)

US 1999-172350P 19991216 (60)

DT Utility

FS APPLICATION

LN.CNT 3036

INCL INCLM: 536/023.500

INCLS: 435/007.210; 435/006.000; 435/325.000; 435/070.210; 435/326.000

NCL NCLM: 536/023.500

NCLS: 435/007.210; 435/006.000; 435/325.000; 435/070.210; 435/326.000

IC [7]

ICM: C12Q001-68

ICS: G01N033-567; C07H021-04; C12P021-04; C12N005-06

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 46 OF 76 USPATFULL on STN

AN 2002:178549 USPATFULL

TI Vaccine for the prevention and treatment of alzheimer's and amyloid related diseases

IN Chalifour, Robert, Ile Bizard, CANADA

Hebert, Lise, Brossard, CANADA

Kong, Xianqi, Dollard-des-Oremaux, CANADA

PI Gervais, Francine, Ile Bizard, CANADA  
US 2002094335 A1 20020718  
AI US 2001-867847 A1 20010529 (9)  
RLI Continuation-in-part of Ser. No. US 2000-724842, filed on 28 Nov 2000,  
PENDING  
PRAI US 1999-168594P 19991129 (60)  
DT Utility  
FS APPLICATION  
LN.CNT 1946  
INCL INCLM: 424/185.100  
NCL NCLM: 424/185.100  
IC [7]  
ICM: A61K039-00  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 47 OF 76 USPATFULL on STN  
AN 2002:99410 USPATFULL  
TI Methods and compositions for the treatment and/or diagnosis of  
neurological diseases and disorders  
IN Solomon, Beka, Herzlia Pituach, ISRAEL  
Frenkel, Dan, Rehovot, ISRAEL  
PI US 2002052311 A1 20020502  
AI US 2001-808037 A1 20010315 (9)  
RLI Continuation-in-part of Ser. No. US 2000-629971, filed on 31 Jul 2000,  
PENDING Continuation-in-part of Ser. No. US 1999-473653, filed on 29 Dec  
1999, PENDING  
PRAI US 1999-152417P 19990903 (60)  
DT Utility  
FS APPLICATION  
LN.CNT 4074  
INCL INCLM: 514/002.000  
INCLS: 424/093.210  
NCL NCLM: 514/002.000  
NCLS: 424/093.210  
IC [7]  
ICM: A61K048-00  
ICS: A61K038-00  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 48 OF 76 USPATFULL on STN  
AN 2002:72861 USPATFULL  
TI Pharmaceutical compositions comprising a soluble laminin receptor  
precursor or a compound which blocks the interaction of the laminin  
receptor precursor and \*\*\*PrPSc\*\*\* or PrPc  
IN Weiss, Stefan, Munich, GERMANY, FEDERAL REPUBLIC OF  
PI US 2002040001 A1 20020404  
AI US 2001-964566 A1 20010928 (9)  
RLI Continuation-in-part of Ser. No. US 2000-424754, filed on 13 Apr 2000,  
PENDING  
PRAI EP 1997-108712 19970530  
WO 1998-EP3220 19980529  
DT Utility  
FS APPLICATION  
LN.CNT 2050  
INCL INCLM: 514/012.000  
INCLS: 435/007.100  
NCL NCLM: 514/012.000  
NCLS: 435/007.100  
IC [7]  
ICM: A61K038-17  
ICS: G01N033-53  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 49 OF 76 USPATFULL on STN  
AN 2002:297296 USPATFULL  
TI Methods for inhibition of membrane fusion-associated events, including  
respiratory syncytial virus transmission  
IN Bolognesi, Dani Paul, Durham, NC, United States  
Matthews, Thomas James, Durham, NC, United States  
Wild, Carl T., Durham, NC, United States  
Barney, Shawn O'Lin, Cary, NC, United States  
Lambert, Dennis Michael, Cary, NC, United States  
Petteway, Stephen Robert, Cary, NC, United States  
Langlois, Alphonse J., Durham, NC, United States  
PA Trimeris, Inc., Durham, NC, United States (U.S. corporation)  
PI US 6479055 B1 20021112

AI US 1995-470896 19950606 (8)  
RLI Continuation-in-part of Ser. No. US 1994-360107, filed on 20 Dec 1994,  
now patented, Pat. No. US 6017536 Continuation-in-part of Ser. No. US  
1994-255208, filed on 7 Jun 1994 Continuation-in-part of Ser. No. US  
1993-73028, filed on 7 Jun 1993, now patented, Pat. No. US 5464933  
DT Utility  
FS GRANTED  
LN.CNT 26553  
INCL INCLM: 424/211.100  
INCLS: 424/186.100; 530/324.000  
NCL NCLM: 424/211.100  
NCLS: 424/186.100; 530/324.000  
IC [7]  
ICM: A61K039-145  
EXF 435/5; 435/240.2; 424/184.1-189.1; 424/204.1-211.1; 424/225.1;  
424/227.1; 424/230.1; 514/1; 514/2; 530/324; 530/350; 530/826  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 50 OF 76 USPATFULL on STN  
AN 2002:81024 USPATFULL  
TI Antibodies specific for native \*\*\*PrPSC\*\*\*  
IN Prusiner, Stanley B., San Francisco, CA, United States  
Williamson, R. Anthony, San Diego, CA, United States  
Burton, Dennis R., La Jolla, CA, United States  
PA The Regents of the University of California, Oakland, CA, United States  
(U.S. corporation)  
The Scripps Research Institute, La Jolla, CA, United States (U.S.  
corporation)  
PI US 6372214 B1 20020416  
AI US 2000-550374 20000413 (9)  
RLI Continuation of Ser. No. US 1998-36579, filed on 6 Mar 1998 Division of  
Ser. No. US 1996-713939, filed on 13 Sep 1996, now patented, Pat. No. US  
5846533, issued on 8 Dec 1998 Continuation-in-part of Ser. No. US  
1995-528104, filed on 14 Sep 1995, now abandoned  
DT Utility  
FS GRANTED  
LN.CNT 2518  
INCL INCLM: 424/130.100  
INCLS: 424/009.100; 424/147.100; 435/007.100; 435/070.100; 435/071.100;  
436/503.000; 436/518.000; 436/547.000; 530/387.100  
NCL NCLM: 424/130.100  
NCLS: 424/009.100; 424/147.100; 435/007.100; 435/070.100; 435/071.100;  
436/503.000; 436/518.000; 436/547.000; 530/387.100  
IC [7]  
ICM: A61K039-395  
ICS: C12P019-00; G01N033-567; G01N033-543; C07K016-00  
EXF 424/9.1; 424/130.1; 424/147.1; 435/7.1; 435/70.1; 435/71.1; 530/387.1;  
436/518; 436/503; 436/547  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 51 OF 76 USPATFULL on STN  
AN 2002:51103 USPATFULL  
TI Methods and compositions for the manufacture of halogenated  
anthracyclines with increased antitumor activity, other anthracyclines,  
halogenated sugars, and glycosyl donors  
IN Priebe, Waldemar, 4239 Emory St., Houston, TX, United States 77005  
Krawczyk, Marta, 175 N. Locust Hill Dr. apt. #2308, Lexington, KY,  
United States 40503  
Skibicki, Piotr, Waszyngton Street 39 Apartment 24, Warsaw 04015, POLAND  
Fokt, Izabela, 1908 Nursery Rd., The Woodlands, TX, United States 77380  
Dziewiszek, Krzysztof, 1908 Nursery Rd., The Woodlands, TX, United  
States 77380  
Grynkiewicz, Grzegorz, .mu.. Zielona 16B/2, 05-092 Lomianki, POLAND  
Perez-Soler, Roman, 564 1st Ave. #20T, New York, NY, United States  
10016  
PI US 6355784 B1 20020312  
AI US 1999-330226 19990610 (9)  
PRAI US 1998-89162P 19980612 (60)  
DT Utility  
FS GRANTED  
LN.CNT 2062  
INCL INCLM: 536/006.400  
INCLS: 536/004.100; 536/017.200; 536/018.400; 536/018.700; 536/122.000  
NCL NCLM: 536/006.400  
NCLS: 536/004.100; 536/017.200; 536/018.400; 536/018.700; 536/122.000  
IC [7]

ICM: C07H015-24

EXF 536/6.4; 536/18.6; 536/122; 536/18.4; 536/18.7; 536/4.1; 536/17.2;  
514/34

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 52 OF 76 SCISEARCH COPYRIGHT (c) 2004 The Thomson Corporation.  
on STN  
AN 2002:986508 SCISEARCH  
GA The Genuine Article (R) Number: 620PU  
TI Induction of antibodies against murine full-length prion protein in  
wild-type mice  
AU Koller M F; Grau T; Christen P (Reprint)  
CS Univ Zurich, Inst Biochem, Winterhurerstr 190, CH-8057 Zurich, Switzerland  
(Reprint); Univ Zurich, Inst Biochem, CH-8057 Zurich, Switzerland  
CYA Switzerland  
SO JOURNAL OF NEUROIMMUNOLOGY, (NOV 2002) Vol. 132, No. 1-2, pp. 113-116.  
Publisher: ELSEVIER SCIENCE BV, PO BOX 211, 1000 AE AMSTERDAM,  
NETHERLANDS.  
ISSN: 0165-5728.  
DT Article; Journal  
LA English  
REC Reference Count: 17  
\*ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS\*

L3 ANSWER 53 OF 76 USPATFULL on STN DUPLICATE 9  
AN 2001:224147 USPATFULL  
TI Prevention of cancer  
IN Henrik, Raskov Hans, Hellerup, Denmark  
PI US 2001049364 A1 20011206  
US 6703380 B2 20040309  
AI US 2001-825891 A1 20010405 (9)  
RLI Continuation-in-part of Ser. No. WO 2000-DK546, filed on 29 Sep 2000,  
UNKNOWN  
PRAI DK 1999-1390 19990929  
DT Utility  
FS APPLICATION  
LN.CNT 1937  
INCL INCLM: 514/164.000  
INCLS: 514/167.000  
NCL NCLM: 514/165.000  
NCLS: 514/167.000  
IC [7]  
ICM: A61K031-616  
ICS: A61K031-59

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 54 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 2001:816487 CAPLUS  
DN 135:356752  
TI Epitope synchronization in antigen presenting cells  
IN Simard, John J. L.; Diamond, David C.; Lei, Xiang-Dong  
PA CTL Immunotherapies Corp., USA  
SO PCT Int. Appl., 131 pp.  
CODEN: PIXXD2  
DT Patent  
LA English  
FAN.CNT 7

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001082963	A2	20011108	WO 2001-US13806	20010427
	WO 2001082963	A3	20020411		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DE, DK, DK, DM, DZ, EE, EE, ES, FI, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	EP 1276896	A2	20030122	EP 2001-930922	20010427
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	JP 2003535824	T2	20031202	JP 2001-579836	20010427
PRAI	US 2000-560465	A	20000428		

US 2000-561074 A 20000428  
US 2000-561571 A 20000428  
US 2000-561572 A 20000428  
WO 2001-US13806 W 20010427

L3 ANSWER 55 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 2001:507867 CAPLUS

DN 135:91527

TI Tissue-specific DNA delivery via M cell-directed vaccines, and enhanced in vivo mucosal IgA and T cell responses resulting therefrom

IN Pascual, David W.

PA Research and Development Institute, Inc., USA

SO PCT Int. Appl., 58 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001049867	A1	20010712	WO 2001-US426	20010108
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	EP 1257654	A1	20021120	EP 2001-901811	20010108
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	US 2004033486	A1	20040219	US 2002-169492	20021021
	US 2004109871	A1	20040610	US 2003-660787	20030912
PRAI	US 2000-174786P	P	20000106		
	WO 2001-US426	W	20010108		
	US 2001-274639P	P	20010312		
	WO 2002-US7254	A2	20020312		
	US 2002-169492	A2	20021021		

RE.CNT 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 56 OF 76 USPATFULL on STN

AN 2001:157792 USPATFULL

TI Antibodies specific for native \*\*\*PrPSc\*\*\*

IN Prusiner, Stanley B., San Francisco, CA, United States

Williamson, R. Anthony, San Diego, CA, United States

Burton, Dennis R., La Jolla, CA, United States

PA The Scripps Research Institute, La Jolla, CA, United States (U.S. corporation)

PI US 6290954 B1 20010918

AI US 1998-36579 19980306 (9)

RLI Division of Ser. No. US 1996-713939, filed on 13 Sep 1996, now patented, Pat. No. US 5846533 Continuation-in-part of Ser. No. US 1995-528104, filed on 14 Sep 1995, now abandoned

DT Utility

FS GRANTED

LN.CNT 2513

INCL INCLM: 424/130.100

INCLS: 424/009.100; 424/147.100; 435/007.100; 435/070.100; 435/071.100; 436/503.000; 436/518.000; 436/547.000; 530/387.100

NCL NCLM: 424/130.100

NCLS: 424/009.100; 424/147.100; 435/007.100; 435/070.100; 435/071.100; 436/503.000; 436/518.000; 436/547.000; 530/387.100

IC [7]

ICM: A61K039-395

ICS: G01N033-53; G01N033-567; C07K016-00

EXF 424/9.1; 424/130.1; 424/147.1; 435/7.1; 435/70.1; 435/71.1; 530/387.1; 436/518; 436/503; 436/547

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 57 OF 76 USPATFULL on STN

AN 2001:125727 USPATFULL

TI Correction of genetic defects using chemical chaperones

IN Welch, William J., San Francisco, CA, United States

Brown, C. Randell, Hershey, PA, United States

PA Tatzelt, Jorg, Munchen, Germany, Federal Republic of  
The Regents of The University of California, Oakland, CA, United states  
(U.S. corporation)  
PI US 6270954 B1 20010807  
AI US 1999-291406 19990413 (9)  
RLI Continuation-in-part of Ser. No. US 1997-838691, filed on 9 Apr 1997,  
now patented, Pat. No. US 5900360  
PRAI US 1996-15155P 19960410 (60)  
DT Utility  
FS GRANTED  
LN.CNT 2342  
INCL INCLM: 435/004.000  
INCLS: 435/026.000; 435/023.000; 435/024.000; 435/963.000  
NCL NCLM: 435/004.000  
NCLS: 435/023.000; 435/024.000; 435/026.000; 435/963.000  
IC [7]  
ICM: C12Q001-00  
ICS: C12Q001-32; C12Q001-37  
EXF 435/4; 435/26; 435/23; 435/24; 435/963  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 58 OF 76 USPATFULL on STN  
AN 2001:67794 USPATFULL  
TI Human respiratory syncytial virus peptides with antifusogenic and  
antiviral activities  
IN Barney, Shawn O'Lin, Cary, NC, United States  
Lambert, Dennis Michael, Cary, NC, United States  
Petteway, Stephen Robert, Cary, NC, United States  
PA Trimeris, Inc., Durham, NC, United States (U.S. corporation)  
PI US 6228983 B1 20010508  
AI US 1995-485264 19950607 (8)  
RLI Division of Ser. No. US 1995-470896, filed on 6 Jun 1995  
Continuation-in-part of Ser. No. US 1994-360107, filed on 20 Dec 1994  
Continuation-in-part of Ser. No. US 1994-255208, filed on 7 Jun 1994  
Continuation-in-part of Ser. No. US 1993-73028, filed on 7 Jun 1993, now  
patented, Pat. No. US 5464933  
DT Utility  
FS Granted  
LN.CNT 32166  
INCL INCLM: 530/300.000  
INCLS: 530/324.000; 530/325.000; 530/326.000; 424/211.100; 424/186.100  
NCL NCLM: 530/300.000  
NCLS: 424/186.100; 424/211.100; 530/324.000; 530/325.000; 530/326.000  
IC [7]  
ICM: A61K038-00  
EXF 530/350; 530/324-329; 530/300; 424/211.1  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 59 OF 76 USPATFULL on STN  
AN 2001:51590 USPATFULL  
TI Clearance and inhibition of conformationally altered proteins  
IN Prusiner, Stanley B., San Francisco, CA, United States  
Supattapone, Surachai, San Francisco, CA, United States  
Scott, Michael, San Francisco, CA, United States  
PA The Regents of the University of California, Oakland, CA, United states  
(U.S. corporation)  
PI US 6214366 B1 20010410  
AI US 1999-322903 19990601 (9)  
DT Utility  
FS Granted  
LN.CNT 1037  
INCL INCLM: 424/405.000  
INCLS: 424/438.000; 424/442.000; 424/484.000; 424/DIG.016; 424/078.320;  
424/078.350; 424/078.360; 424/078.370; 424/078.380; 514/772.300;  
514/772.400; 514/772.500; 514/772.600; 514/772.700  
NCL NCLM: 424/405.000  
NCLS: 424/078.320; 424/078.350; 424/078.360; 424/078.370; 424/078.380;  
424/438.000; 424/442.000; 424/484.000; 424/DIG.016; 514/772.300;  
514/772.400; 514/772.500; 514/772.600; 514/772.700  
IC [7]  
ICM: A01N025-10  
EXF 424/78.32; 424/78.35; 424/78.38; 424/405; 424/438; 424/442; 424/DIG.16;  
514/772.3-772.7  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 60 OF 76 WPIDS COPYRIGHT 2004 THOMSON DERWENT on STN

AN 2002-049350 [06] WPIDS  
DNN N2002-036483 DNC C2002-013896  
TI New polypeptides, useful as antiviral agents, comprise their prion proteins able to bind nucleic acid, nucleocapsid proteins, and ligands for use as antiprion agents.  
DC B04 C06 D16 S03  
IN DARLIX, J L; GABUS, D C; LEBLANC, P; DARLIX, J; GABUS-DARLIX, C  
PA (INRM) INSERM INST NAT SANTE & RECH MEDICALE  
CYC 95  
PI WO 2001083747 A2 20011108 (200206)\* FR 80 C12N015-12  
RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW MZ  
NL OA PT SD SE SL SZ TR TZ UG ZW  
W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK  
DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ  
LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ PL PT RO RU SD  
SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW  
FR 2808278 A1 20011102 (200206) C07K014-435  
AU 2001056454 A 20011112 (200222) C12N015-12  
ADT WO 2001083747 A2 WO 2001-FR1336 20010430; FR 2808278 A1 FR 2000-5535  
20000428; AU 2001056454 A AU 2001-56454 20010430  
FDT AU 2001056454 A Based on WO 2001083747  
PRAI FR 2000-5535 20000428  
IC ICM C07K014-435; C12N015-12  
ICS A61K038-17; A61K039-395; A61K048-00; A61P025-00; A61P031-12;  
C07K014-47; C07K019-00; C12N007-00; C12N015-86; C12Q001-68;  
C12Q001-70; G01N033-569; G01N033-68

L3 ANSWER 61 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED.  
on STN DUPLICATE 10

AN 2001435095 EMBASE  
TI Human aqueous humor levels of oral ciprofloxacin, levofloxacin, and moxifloxacin.  
AU Garcia-Saenz M.C.; Arias-Puente A.; Fresnadillo-Martinez M.J.; Carrasco-Font C.  
CS Dr. M.C. Garcia-Saenz, Fundacion Hospital de Alcorcon, Budapest 1, 28922 Alcorcon, Madrid, Spain. mcgarcias@fhalcorcon.es  
SO Journal of Cataract and Refractive Surgery, (2001) 27/12 (1969-1974).  
Refs: 23  
ISSN: 0886-3350 CODEN: JCSUEV  
PUI S 0886-3350(01)00997-X  
CY United States  
DT Journal; Article  
FS 012 Ophthalmology  
037 Drug Literature Index  
LA English  
SL English

L3 ANSWER 62 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED.  
on STN

AN 2003330992 EMBASE  
TI Short-term effect of mitomycin-C augmented trabeculectomy on axial length and corneal astigmatism.  
AU Kook M.S.; Kim H.B.; Lee S.U.  
CS Dr. M.S. Kook, Department of ophthalmology, Ulsan University School of Medicine, Asan Medical Center, 388-1 Pungnap-dong, Songpa-gu, Seoul, 138-736, Korea, Republic of  
SO Journal of Cataract and Refractive Surgery, (2001) 27/4 (518-523).  
Refs: 25  
ISSN: 0886-3350 CODEN: JCSUEV  
PUI S 0886-3350(00)00646-5  
CY United States  
DT Journal; Article  
FS 012 Ophthalmology  
037 Drug Literature Index  
038 Adverse Reactions Titles  
LA English  
SL English

L3 ANSWER 63 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN

AN 2000:861780 CAPLUS  
DN 134:28436  
TI Vaccines against conformation-dependent protein and non-protein antigens  
IN Goletz, Steffen; Karsten, Uwe  
PA Max-Delbrück-Centrum fuer Molekulare Medizin, Germany  
SO PCT Int. Appl., 36 pp.  
CODEN: PIXXD2

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000073430	A2	20001207	WO 2000-DE1809	20000529
	WO 2000073430	A3	20010329		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	DE 10027695	A1	20010419	DE 2000-10027695	20000529
	EP 1181058	A2	20020227	EP 2000-951201	20000529
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
PRAI	DE 1999-19924405	A	19990527		
	DE 1999-19943016	A	19990909		
	WO 2000-DE1809	W	20000529		

L3 ANSWER 64 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN

AN 2000:529259 CAPLUS

DN 133:132106

TI Prion gene knockout animal cell lines, antiviral vaccine production, and mutant prion gene detection in mammals

IN Yokoyama, Takashi; Itohara, Shigemi; Onodera, Takashi

PA Ministry of Agriculture, Forestry and Fisheries National Institute of Animal, Japan

SO Jpn. Kokai Tokkyo Koho, 10 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 2000210078	A2	20000802	JP 1999-13834	19990122
	JP 3084399	B2	20000904		
PRAI	JP 1999-13834		19990122		

L3 ANSWER 65 OF 76 WPIDS COPYRIGHT 2004 THOMSON DERWENT on STN

AN 2001-070921 [08] WPIDS

CR 2004-411388 [38]

DNC C2001-019767

TI Pharmaceutical composition comprising immunogen against amyloid component such as fibril peptide or protein, or antibody against amyloid component useful for treating amyloid diseases or amyloidoses.

DC B04 D16 P31

IN SCHENK, D B; MASLIAH, E

PA (NEUR-N) NEURALAB LTD; (MASL-I) MASLIAH E; (SCHE-I) SCHENK D B

CYC 94

PI	WO 2000072876	A2	20001207 (200108)*	EN 140	A61K039-00
	RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW MZ				
	NL OA PT SD SE SL SZ TZ UG ZW				
	W: AE AG AL AM AT AU AZ BA BB BG BR BY CA CH CN CR CU CZ DE DK DM DZ				
	EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC				
	LR LS LT LU LV MA MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI				
	SK SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW				
	AU 2000053163	A	20001218 (200118)		
	NO 2001005758	A	20020130 (200223)		A61K000-00
	EP 1185296	A2	20020313 (200225)	EN	A61K039-00
	R: AL AT BE CH CY DE DK ES FI FR GB GR IE IT LI LT LU LV MC MK NL PT				
	RO SE SI				
	BR 2000011103	A	20020319 (200228)		A61K039-00
	HU 2002001205	A2	20020828 (200264)		A61K039-00
	KR 2002025884	A	20020404 (200266)		A61K038-00
	SK 2001001718	A3	20020910 (200274)		A61K039-00
	CZ 2001004154	A3	20021113 (200282)		A61K039-00
	CN 1377278	A	20021030 (200314)		A61K039-00
	JP 2003516929	W	20030520 (200334)	166	A61K045-08
	ZA 2001009662	A	20030730 (200355)	155	A61K000-00
	MX 2001012293	A1	20021201 (200377)		A61K039-00
	US 2004146521	A1	20040729 (200450)		A61K039-00

ADT WO 2000072876 A2 WO 2000-US15239 20000601; AU 2000053163 A AU 2000-53163  
20000601; NO 2001005758 A WO 2000-US15239 20000601, NO 2001-5758 20011126;  
EP 1185296 A2 EP 2000-938075 20000601, WO 2000-US15239 20000601; BR  
2000011103 A BR 2000-11103 20000601, WO 2000-US15239 20000601; HU  
2002001205 A2 WO 2000-US15239 20000601, HU 2002-1205 20000601; KR  
2002025884 A KR 2001-715508 20011201; SK 2001001718 A3 WO 2000-US15239  
20000601, SK 2001-1718 20000601; CZ 2001004154 A3 WO 2000-US15239  
20000601, CZ 2001-4154 20000601; CN 1377278 A CN 2000-808358 20000601; JP  
2003516929 W WO 2000-US15239 20000601, JP 2001-511318 20000601; ZA  
2001009662 A ZA 2001-9662 20011123; MX 2001012293 A1 WO 2000-US15239  
20000601, MX 2001-12293 20011129; US 2004146521 A1 Provisional US  
1999-137010P 19990601, CIP of US 2000-580015 20000526, CIP of US  
2000-585817 20000601, Provisional US 2002-423012P 20021101, US 2003-698099  
20031031

FDT AU 2000053163 A Based on WO 2000072876; EP 1185296 A2 Based on WO  
2000072876; BR 2000011103 A Based on WO 2000072876; HU 2002001205 A2 Based  
on WO 2000072876; SK 2001001718 A3 Based on WO 2000072876; CZ 2001004154  
A3 Based on WO 2000072876; JP 2003516929 W Based on WO 2000072876; MX  
2001012293 A1 Based on WO 2000072876

PRAI US 1999-137010P 19990601; US 2000-580015 20000526;  
US 2000-585817 20000601; US 2002-423012P 20021101;  
US 2003-698099 20031031

IC ICM A61K000-00; A61K038-00; A61K039-00; A61K045-08  
ICS A61K039-385; A61K039-39; A61K039-395; A61K039-44; A61K047-48;  
A61K048-00; A61P001-04; A61P003-00; A61P017-00; A61P017-06;  
A61P019-00; A61P019-02; A61P025-28; A61P029-00; A61P035-00;  
A61P037-00; A61P043-00; G01N033-68

ICA C12N015-09

L3 ANSWER 66 OF 76 Elsevier BIOBASE COPYRIGHT 2004 Elsevier Science B.V.  
on STN DUPLICATE

AN 2000253205 ESBIOBASE

TI Successful treatment of metastatic retinoblastoma

AU Dunkel I.J.; Aledo A.; Kernan N.A.; Kushner B.; Bayer L.; Gollamudi S.V.;  
Finlay J.L.; Abramson D.H.

CS Dr. I.J. Dunkel, Department of Pediatrics, Mem. Sloan-Kettering Cancer  
Center, Box 185, 1275 York Avenue, New York, NY 10021, United States.  
E-mail: dunkeli@mskcc.org

SO Cancer, (15 NOV 2000), 89/10 (2117-2121), 17 reference(s)

CODEN: CANCAR ISSN: 0008-543X

DT Journal; Article

CY United States

LA English

SL English

L3 ANSWER 67 OF 76 USPATFULL on STN

AN 1999:53572 USPATFULL

TI Correction of genetic defects using chemical chaperones

IN Welch, William J., 48 Fountain, San Francisco, CA, United States 94114  
Brown, C. Randell, 1470 9th Ave. #12, San Francisco, CA, United States  
94122

PI Tatzelt, Jorg, 740 Parnassus, San Francisco, CA, United States 94122  
US 5900360 19990504

AI US 1997-838691 19970409 (8)

PRAI US 1996-15155P 19960410 (60)

DT Utility

FS Granted

LN.CNT 2062

INCL INCLM: 435/029.000

INCLS: 435/004.000; 435/005.000; 435/034.000; 436/063.000; 436/086.000;  
436/506.000; 436/811.000

NCL NCLM: 435/029.000

NCLS: 435/004.000; 435/005.000; 435/034.000; 436/063.000; 436/086.000;  
436/506.000; 436/811.000

IC [6]

ICM: C12Q001-02

EXF ICS: C12Q001-04; G01N033-48; G01N033-564

435/29; 435/34; 435/4; 435/5; 436/86; 436/63; 436/811; 436/506

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 68 OF 76 DRUGU COPYRIGHT 2004 THOMSON DERWENT on STN DUPLICATE  
12

AN 1998-11346 DRUGU T S

TI Much ado about not... enough data: high-dose chemotherapy with autologous  
stem cell rescue for breast cancer.

AU Zujewski J; Nelson A; Abrams J

LO Bethesda; Potomac, Md., USA  
SO J.Natl.Cancer Inst. (90, No. 3, 200-91, 1998) 1 Fig. 8 Tab. 51 Ref.  
CODEN: JNCIEQ ISSN: 0027-8874  
AV National Institutes of Health, Bldg. 10 Rm. 12N/226, Bethesda, MD 20892,  
U.S.A.  
LA English  
DT Journal  
FA AB; LA; CT  
FS Literature

L3 ANSWER 69 OF 76 PASCAL COPYRIGHT 2004 INIST-CNRS. ALL RIGHTS RESERVED.  
on STN  
AN 1996-0380308 PASCAL  
CP Copyright .COPYRGT. 1996 INIST-CNRS. All rights reserved.  
TIEN High dose consolidation with autologous stem cell rescue ( \*\*\*ASCR\*\*\* )  
for nephroblastoma initially treated according to the SIOP 9/GPOH trial  
and study  
AU HEMPEL L.; KREMENS B.; WEIRICH A.; GRAF N.; ZINTL F.; LUDWIG R.  
CS Dept. of Pediatric Hematology and Oncology, University of Jena, Germany,  
Federal Republic of  
SO Klinische Paediatrie, (1996), 208(4), 186-189, 18 refs.  
ISSN: 0300-8630 CODEN: KLPDB2  
DT Journal  
BL Analytic  
CY Germany, Federal Republic of  
LA English  
SL German  
AV INIST-4105, 354000060429320080

L3 ANSWER 70 OF 76 CANCERLIT on STN  
AN 96604645 CANCERLIT  
DN 96604645  
TI High-dose chemotherapy with bone marrow transplantation in the treatment  
of breast cancer (Meeting abstract).  
AU Fields K K; Elfenbein G J  
CS Div. of Bone Marrow Transplantation, Univ. of South Florida, Tampa, FL  
33612.  
SO Cancer Invest, (1995) 13 (Suppl 1) 12-3.  
ISSN: 0735-7907.  
DT (MEETING ABSTRACTS)  
LA English  
FS Institute for Cell and Developmental Biology  
EM 199605  
ED Entered STN: 19970509  
Last Updated on STN: 19970509

L3 ANSWER 71 OF 76 CABA COPYRIGHT 2004 CABI on STN DUPLICATE 13  
AN 96:29330 CABA  
DN 19960101143  
TI Ablation of the prion protein (PrP) gene in mice prevents scrapie and  
facilitates production of anti-PrP antibodies  
AU Prusiner, S. B.; Groth, D.; Serban, A.; Koehler, R.; Foster, D.; Torchia,  
M.; Burton, D.; Yang ShuLian; DeArmond, S. J.; Yang, S. L.  
CS Department of Neurology, University of California, San Francisco, CA  
94143, USA.  
SO Proceedings of the National Academy of Sciences of the United States of  
America, (1993) Vol. 90, No. 22, pp. 10608-10612. 53 ref.  
ISSN: 0027-8424  
DT Journal  
LA English  
ED Entered STN: 19960318  
Last Updated on STN: 19960318

L3 ANSWER 72 OF 76 CANCERLIT on STN  
AN 95606623 CANCERLIT  
DN 95606623  
TI High dose mitoxantrone (M) thiotapec (T) and cyclophosphamide (C) plus  
autologous stem cell rescue ( \*\*\*ASCR\*\*\* ) in patients with breast  
cancer (Meeting abstract).  
AU Taylor C W; List A F; Azar C A; Rifkin R M; Mosley K; Dalton W S  
CS Arizona Cancer Center, Tucson AZ 85724.  
SO Breast Cancer Treat Res, (1993) 27 (1/2) 183.  
ISSN: 0167-6806.  
DT Journal; Article; (JOURNAL ARTICLE)  
LA English  
FS Institute for Cell and Developmental Biology

EM 199503  
ED Entered STN: 19950313  
Last Updated on STN: 19950313

L3 ANSWER 73 OF 76 CANCERLIT on STN DUPLICATE 14  
AN 94073488 CANCERLIT  
DN 94073488 PubMed ID: 7902764  
TI Intensive dose ifosfamide, carboplatin, and etoposide followed by autologous stem cell rescue: results of a phase I/II study in breast cancer patients.  
AU Fields K K; Perkins J P; Hiemenz J W; Zorsky P E; Janssen W E; Kronish L E; Machak M C; Elfenbein G J  
CS Department of Internal Medicine, H. Lee Moffitt Cancer Center, University of South Florida, Tampa 33612.  
SO SURGICAL ONCOLOGY, (1993) 2 (1) 87-95.  
Journal code: 9208188. ISSN: 0960-7404.  
CY ENGLAND: United Kingdom  
DT (CLINICAL TRIAL)  
(CLINICAL TRIAL, PHASE I)  
(CLINICAL TRIAL, PHASE II)  
LA Journal; Article; (JOURNAL ARTICLE)  
English  
FS MEDLINE; Priority Journals  
OS MEDLINE 94073488  
EM 199401  
ED Entered STN: 19941107  
Last Updated on STN: 19950508

L3 ANSWER 74 OF 76 DRUGU COPYRIGHT 2004 THOMSON DERWENT on STN  
AN 1992-26777 DRUGU T S  
TI Ifosfamide, Carboplatin and Etoposide (ICE) with Autologous Stem Cell Rescue ( \*\*\*ASCR\*\*\* ): Toxicities.  
AU Elfenbein G; Fields K; Zorsky P; Hiemenz J; Janssen W; Perkins J  
LO Tampa, Florida, United States  
SO Proc.Am.Soc.Clin.Oncol. (11, 28 Meet., 90, 1992)  
AV Div. Bone Marrow Transplant, Univ. of South Fla., Tampa, FL, U.S.A. (11 authors).  
LA English  
DT Journal  
FA AB; LA; CT  
FS Literature

L3 ANSWER 75 OF 76 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on STN DUPLICATE 15  
AN 1990:91415 BIOSIS  
DN PREV199089050766; BA89:50766  
TI HIGH-DOSE CONSOLIDATION THERAPY WITH AUTOLOGOUS STEM CELL RESCUE IN STAGE IV BREAST CANCER.  
AU WILLIAMS S F [Reprint author]; MICK R; DESSER R; GOLICK J; BESCHORNER J; BITRAN J D  
CS UNIV CHICAGO, BOX 420, 5841 S MARYLAND AVE, CHICAGO, ILL 60637, USA  
SO Journal of Clinical Oncology, (1989) Vol. 7, No. 12, pp. 1824-1830.  
CODEN: JCONDN. ISSN: 0732-183X.  
DT Article  
FS BA  
LA ENGLISH  
ED Entered STN: 9 Feb 1990  
Last Updated on STN: 9 Feb 1990

L3 ANSWER 76 OF 76 DRUGU COPYRIGHT 2004 THOMSON DERWENT on STN  
AN 1990-16751 DRUGU T S  
TI A Phase II Study of Induction Chemotherapy Followed by Intensification with High Dose Chemotherapy with Autologous Stem Cell Rescue ( \*\*\*ASCR\*\*\* ) in Stage IV Breast Cancer.  
AU Williams S; Bitran J; Desser R; Golick J; Beschorner J; Fullem L  
LO Chicago, Illinois, United States  
SO Proc.Am.Soc.Clin.Oncol. (7, 24 Meet., 9, 1988)  
AV Joint Section of Hematology/Oncology, University of Chicago and Michael Reese Medical Centers, Chicago, IL 60637, U.S.A.  
LA English  
DT Journal  
FA AB; LA; CT  
FS Literature

STN INTERNATIONAL LOGOFF AT 17:07:16 ON 09 SEP 2004